



THE UNIVERSITY *of* EDINBURGH

Edinburgh Research Explorer

A Tgfb β 1/Snai1-dependent developmental module at the core of vertebrate axial elongation

Citation for published version:

Dias, A, Lozovska, A, Wymeersch, FJ, Nóvoa, A, Binagui-Casas, A, Sobral, D, Martins, GG, Wilson, V & Mallo, M 2020, 'A Tgfb β 1/Snai1-dependent developmental module at the core of vertebrate axial elongation', *eLIFE*, vol. 9. <https://doi.org/10.7554/eLife.56615>

Digital Object Identifier (DOI):

[10.7554/eLife.56615](https://doi.org/10.7554/eLife.56615)

Link:

[Link to publication record in Edinburgh Research Explorer](#)

Document Version:

Publisher's PDF, also known as Version of record

Published In:

eLIFE

General rights

Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy

The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.



A Tgfb β 1/Snai1-dependent developmental module at the core of vertebrate axial elongation

André Dias¹, Anastasiia Lozovska^{1†}, Filip J Wymeersch^{2†‡}, Ana Nóvoa¹, Anahi Binagui-Casas², Daniel Sobral^{1§}, Gabriel G Martins^{1,3}, Valerie Wilson², Moises Mallo^{1*}

¹Instituto Gulbenkian de Ciência, Oeiras, Portugal; ²Centre for Regenerative Medicine, Institute for Regeneration and Repair, University of Edinburgh, Edinburgh, United Kingdom; ³Faculdade de Ciências da Universidade de Lisboa, Lisboa, Portugal

*For correspondence:
mallo@igc.gulbenkian.pt

†These authors contributed
equally to this work

Present address: [‡]RIKEN Center
for Biosystems Dynamics
Research, Hyogo, Japan;
[§]UCIBIO, Departamento de
Ciências da Vida, Faculdade de
Ciências e Tecnologia,
Universidade NOVA de Lisboa,
Caparica, Portugal

Competing interests: The
authors declare that no
competing interests exist.

Funding: See page 23

Received: 03 March 2020

Accepted: 18 June 2020

Published: 29 June 2020

Reviewing editor: Elizabeth
Robertson, University of Oxford,
United Kingdom

© Copyright Dias et al. This
article is distributed under the
terms of the [Creative Commons
Attribution License](#), which
permits unrestricted use and
redistribution provided that the
original author and source are
credited.

Abstract Formation of the vertebrate postcranial body axis follows two sequential but distinct phases. The first phase generates pre-sacral structures (the so-called primary body) through the activity of the primitive streak on axial progenitors within the epiblast. The embryo then switches to generate the secondary body (post-sacral structures), which depends on axial progenitors in the tail bud. Here we show that the mammalian tail bud is generated through an independent functional developmental module, concurrent but functionally different from that generating the primary body. This module is triggered by convergent Tgfb β 1 and Snai1 activities that promote an incomplete epithelial to mesenchymal transition on a subset of epiblast axial progenitors. This EMT is functionally different from that coordinated by the primitive streak, as it does not lead to mesodermal differentiation but brings axial progenitors into a transitory state, keeping their progenitor activity to drive further axial body extension.

Introduction

Formation of the vertebrate body is a complex and dynamic process involving a series of sequential growth and patterning activities. In amniotes, the primordia of the different organs and body structures are laid down progressively in a head to tail sequence by dedicated axial progenitors with stem cell-like properties (*Stern et al., 2006*). Although continuous, this process can be divided into distinct steps based on structural and regulatory features. At the end of gastrulation, during the so-called primary body formation (i.e. post-occipital region of the head, neck and trunk) axial progenitors are located in the epiblast, an epithelial layer at the caudal embryonic end, and their activity is organized by the primitive streak (PS) (*Aires et al., 2018; Holmdahl, 1925; Steventon and Martinez Arias, 2017*). At this stage, the embryo contains axial progenitors with different potencies. These include the neuro-mesodermal progenitors (NMPs), a bipotent cell population that can generate both neural and mesodermal tissues, and the lateral and paraxial mesoderm progenitors (LPMPs), with potential limited to mesodermal lineages, that together with the endoderm generate organic systems involved in most vital and reproductive functions (*Wilson et al., 2009; Wymeersch et al., 2016*). After PS regression and caudal neuropore closure, the embryo engages in secondary body formation (essentially the tail). At this stage, axial progenitors, which are now restricted to NMPs, are located in the tail bud (*Bénazéraf and Pourquié, 2013; Henrique et al., 2015; Steventon and Martinez Arias, 2017; Wilson et al., 2009*).

A large number of genetic experiments led to the identification of factors regulating axial progenitor activity. Some of these factors, including *Wnt3a*, *Fgf8*, *T(Brachyury)* or the *Cdx* gene family,

are required during both primary and secondary body axis formation, as their partial or total inactivation produce different degrees of axial truncations depending on the levels of gene activity left available to the axial progenitors (Boulet and Capecchi, 2012; Greco et al., 1996; Herrmann et al., 1990; Naiche et al., 2011; Savory et al., 2011; Takada et al., 1994). Other factors show regional specific activity, determining whether progenitors generate trunk or tail structures (Aires et al., 2019; Aires et al., 2018; Aires et al., 2016; DeVeale et al., 2013; Robinton et al., 2019; Wymeersch et al., 2019). Gain and loss of function experiments in the mouse revealed a central role for *Pou5f1* (also known as *Oct4*) in trunk development. Indeed, conditional *Pou5f1* inactivation after it had fulfilled its role during preimplantation and early post-implantation stages resulted in embryos lacking trunk structures but still containing recognizable tails (DeVeale et al., 2013). Conversely, sustained transgenic *Pou5f1* expression in the axial progenitor region extended trunk development at the expense of the tail (Aires et al., 2016). *Pou5f1* importance for vertebrate trunk development was further revealed by the finding that the remarkably long trunks characteristic of the snake body plan seemed to derive from a chromosomal rearrangement involving the *Pou5f1* locus during vertebrate evolution that placed this gene under the control of regulatory elements that maintained its expression for very long developmental periods (Aires et al., 2016).

In the tail bud, axial progenitor activity is independent of *Pou5f1* (DeVeale et al., 2013). Genetic experiments in mouse embryos revealed that in this area the *Lin28/let-7* axis together with *Hox13* genes, particularly those belonging to the *HoxB* and *HoxC* clusters, occupy a prevalent position in the regulatory hierarchy of axial progenitors in the tail bud (Aires et al., 2019; Robinton et al., 2019). Interestingly, while tail bud progenitors show drastic responses to variations in the *Lin28/let-7* pathway and are strongly inhibited by premature activation of *Hox13* genes, their trunk counterparts are largely non-responsive to those activities (Aires et al., 2019; Robinton et al., 2019), suggesting the existence of differences in cell competence, at the progenitor level, during primary and secondary body formation. The differences in progenitor regulation at trunk and tail levels seem to be also associated with changes in their functional characteristics. For instance, while NMPs produce neural tube at both axial levels, they are thought to follow different mechanisms in the two regions (Catala et al., 1995; Schoenwolf, 1984; Schoenwolf and Smith, 1990). Differences can also be observed in the properties of their mesodermal derivatives. In particular, while disconnecting the *Lfng* cycling activity disturbs somitogenesis at trunk levels, it has no or minor effects in the tail (Shifley et al., 2008; Williams et al., 2014). Conversely, forced *Hoxb6* expression blocks tail somitogenesis but has no effect at trunk levels (Casaca et al., 2016).

Despite these major differences in axial progenitor regulation and competence, lineage tracing experiments indicate that post-occipital neural and mesodermal structures are generated from a progenitor pool that is, at least to some extent, maintained along the main vertebrate body axis (Tzouanacou et al., 2009), thus implying that transition from primary to secondary body development entails unknown molecular mechanisms occurring at the progenitor level (Aires et al., 2018). We have previously shown that *Gdf11* activity plays a relevant role in this process (Aires et al., 2019; Jurberg et al., 2013). However, partial redundancy by *Gdf8* (McPherron et al., 2009) complicates proper evaluation of this process, as trunk-to-tail transition eventually becomes activated in *Gdf11* mutant embryos, although at more caudal axial levels and generating a number of abnormal structures (Aires et al., 2016; McPherron et al., 1999). Therefore, to evaluate the underlying molecular mechanisms, we compared the molecular characteristics of axial progenitors at progressively later developmental stages using a single-cell RNA-sequencing (scRNA-seq) approach and found that the switch from primary to secondary body development entails an incomplete epithelial to mesenchymal transition (EMT) affecting a subset of axial progenitors in the epiblast. Further analyses showed that this process is functionally different from that generating primary body structures and that it is driven by the sequential activity of *Tgfb β 1* and *Snai1*. Together, our data uncovered a distinct functional developmental module generating the tail bud from a subset of epiblast axial progenitors, that initiates secondary body formation in mouse embryos.

Results

Axial progenitors undergo an incomplete EMT during axial extension

To evaluate the underlying mechanisms of the changes in progenitor activity as they switch from generating primary to secondary body structures, we compared their molecular characteristics at progressively later developmental stages using a scRNA-seq approach. We first obtained scRNA-seq data from the caudal lateral epiblast (CLE) of early head fold mouse embryos [~embryonic stage (E) 8.0], which contain axial progenitors (Cambray and Wilson, 2002; Tam and Behringer, 1997; Figure 1A). The dissected tissue included some nascent mesoderm, but these cells segregated from those of the epiblast using the single-cell consensus clustering (SC3) framework (Kiselev et al., 2017), producing two well-defined clusters (Figure 1A and Figure 1—figure supplement 1A). We then compared these data with published scRNA-seq data from CLE regions of E8.5 and E9.5 embryos (Gouti et al., 2017), which similarly to the E8.0 single-cells, contained some nascent mesoderm. Principal component analysis (PCA) of all these single-cell transcriptomes indicated that the epiblast cluster from E8.0 embryos becomes more similar to the mesodermal component (Figure 1A; Figure 1—figure supplement 1B,C). Interestingly, differential gene expression analysis revealed an increase of mesenchyme-associated genes and a concomitant decrease of some epithelial markers (Kalluri and Weinberg, 2009; Lamouille et al., 2014) in epiblast clusters as development proceeds (Figure 1B). For instance, we observed a transition from *Cdh1* (*E-cadherin*) to *Cdh2* (*N-cadherin*), as well as downregulation of *Epcam* and up-regulation of *Vim*. This tendency was also observed in cells co-expressing *Sox2* and *T* (Figure 2), typically associated with NMPs (Cambray and Wilson, 2007; Koch et al., 2017; Martin and Kimelman, 2012; Olivera-Martinez et al., 2012; Tsakiridis et al., 2014; Wymeersch et al., 2016). These results suggest that progress into tail bud stages requires that axial progenitors undergo an EMT.

We further tested this idea in two additional ways. As the majority of axial progenitors in the Chordoneural Hinge (CNH; the region of the tail bud that drives tail elongation) are descended from cells in the Node-Streak Border (NSB) (Cambray and Wilson, 2007), we compared the transcriptome of these two regions of the mouse embryo using data from Wymeersch et al., 2019. This analysis revealed increased expression of mesenchymal-related genes (e.g. *Snai1*, *Vim* and *Zeb1*) and downregulation of some epithelial markers (e.g. *Cdh1*, *Krt18* and *Cldn4*) in the CNH (Figure 1D). In a complementary approach, we examined the transcriptome of a cell population highly enriched in axial progenitors isolated from E10.5 tail buds (de Lemos et al., 2019), using as a reference the transcriptome of unsorted tail bud tissue (Aires et al., 2019). Again, many mesenchymal markers were highly expressed in the sorted axial progenitors, at levels comparable to those observed in the total tail bud (Figure 1E). Conversely, expression of many epithelial markers was significantly lower in the progenitor pool than in the reference tail bud. Therefore, the gene expression profile of tail bud axial progenitors is more consistent with a mesenchymal than with an epithelial phenotype, further suggesting that tail bud formation entails an EMT on the axial progenitors. Interestingly, however, in all our analyses the loss of epithelial characteristics in tail bud axial progenitors was not complete. Indeed, some epithelial markers maintained high expression levels (e.g. *Tjp1*, *Lamb1*) or were upregulated (e.g. *Krt10* in the CNH) when axial progenitors moved into the tail bud (Figure 1D,E), thus indicating that the EMT involved in this process could be classified into the category of incomplete or transitional EMTs (further referred as tb-EMT), more akin to those described in metastatic processes than to those driving embryonic development (Kalluri and Weinberg, 2009; Lamouille et al., 2014; Nieto et al., 2016; Pastushenko et al., 2018). Interestingly, while during gastrulation axial progenitors undergoing EMT through the PS enter differentiation routes (e.g. mesoderm formation), tb-EMT keeps the progenitor properties of these cells, capacitating them to drive further axial extension from the tail bud (Aires et al., 2019; Rodrigo Albors et al., 2018; Tzouanacou et al., 2009). This further indicates that although gastrulation and tail bud formation both involve EMTs on axial progenitors within the epiblast, they seem to be functionally separate processes, suggesting that they should rely on different regulatory mechanisms.

Snai1 is required for axial progenitor mobilization to form the tail bud

Snail genes are among the most prominent EMT regulators in physiological and pathological processes (Barrallo-Gimeno and Nieto, 2005; Carver et al., 2001; Lomeli et al., 2009; Murray and

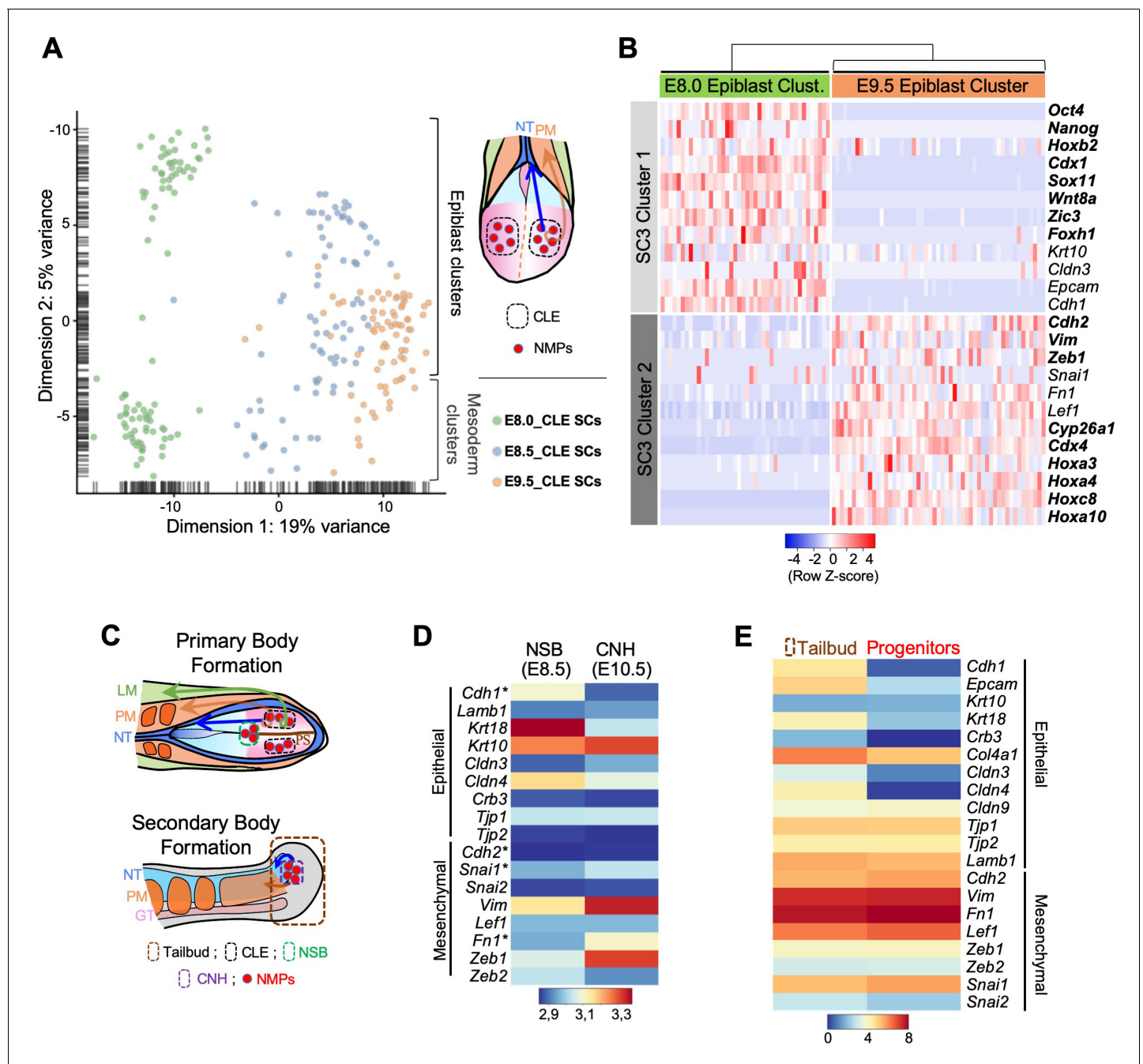


Figure 1. Axial progenitors undergo an incomplete EMT during axial extension. (A) PCA analysis of CLE scRNA-seq datasets from E8.0, E8.5 and E9.5 embryos. Dimension one represents developmental time, whereas dimension two shows differences in cell characteristics. Epiblast and mesodermal clusters are indicated. During primary body formation, epiblast clusters converge towards the mesodermal compartment. (B) Z-score heatmap gene expression analysis of selected genes and marker genes (highlighted in bold), obtained with the SC3 when pre-clustered E8.0 and E9.5 epiblast single-cells are forced to form two clusters. Downregulation of some epithelial-associated genes (e.g. *Cdh1* and *Epcam*) together with upregulation of several mesenchymal-related genes (e.g. *Cdh2*, *Vim*, *Zeb1*) at E9.5 is observed. P-values are shown in **Figure 1—source data 1**. (C) Diagrams showing the regions used for the analysis in D and E. (D) Comparative heatmap representation of microarray gene expression between E8.5 node-streak border (NSB) and E10.5 chordoneural hinge (CNH). Epithelial and mesenchymal markers are indicated. (E) Comparative heatmap representation of RNA-seq gene expression in tail bud axial progenitors and a similar-staged reference total tail bud. Both (D and E) analysis indicated that loss of epithelial markers in tail bud axial progenitors is not complete (e.g. *Tjp1* and *Krt10*). NT: Neural Tube; PM: Paraxial Mesoderm; LM: Lateral Mesoderm; GT: Gut; PS: Primitive Streak; * means Average.

The online version of this article includes the following source data and figure supplement(s) for figure 1:

Source data 1. p-values corresponding to the SC3 analysis of RNA-seq values represented in **Figure 1** and **Figure 1—figure supplement 1**.

Figure 1 continued on next page

Figure 1 continued

Figure supplement 1. SC3 clustering analysis enables the distinction of epiblast from early mesoderm cells.

Gridley, 2006; Nieto, 2002; Zeisberg and Neilson, 2009). From this gene family, *Snai1* is a good candidate to be involved in EMT processes associated with axial extension, as in mouse embryos it is expressed in the PS region starting at early developmental stages (Hernández-Martínez et al., 2019; Nieto et al., 1992; Figure 2—figure supplement 1). Indeed, gene inactivation experiments in mouse indicated that *Snai1* is involved in EMT processes during gastrulation (Carver et al., 2001).

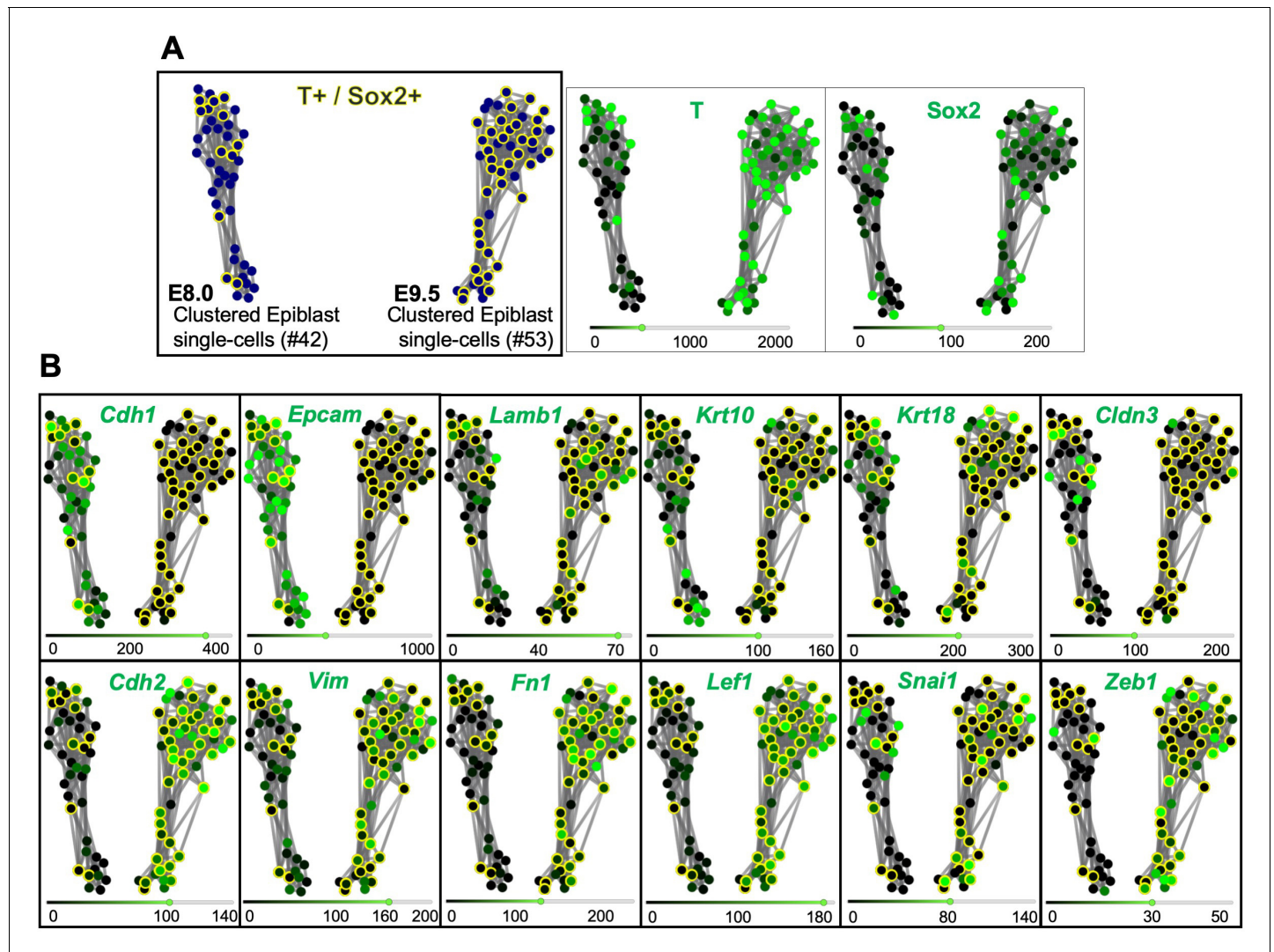


Figure 2. *T* and *Sox2* double-positive cells acquire mesenchymal properties during primary body formation. (A) RPKM (Reads per kilo base per million mapped reads) SPRING analysis of pre-clustered epiblast E8.0 and E9.5 single cells, organizes them in two clusters according to their developmental stage. *T* and *Sox2* double-positive cells are shown circled in yellow. (B) Comparative gene expression analysis of epithelial (upper row) and mesenchymal (lower row) markers focused on *T*⁺ and *Sox2*⁺ epiblast cells suggest that they undergo an EMT between E8.0 and E9.5. The scRNA-seq values in RPKM used for this analysis are shown in Figure 2—source data 1.

The online version of this article includes the following source data and figure supplement(s) for figure 2:

Source data 1. RPKM (Reads per kilo base per million mapped reads) values represented in Figure 2, Figure 2—figure supplement 1 and Figure 2—figure supplement 2.

Figure supplement 1. *Snai1* is expressed at low levels in regions known to contain axial progenitors.

Figure supplement 2. *Tgfb1* is expressed in regions known to contain axial progenitors.

Interestingly, when this early *Snai1* function was bypassed through a conditional approach that inactivated this gene in the epiblast using the *Meox2-Cre* driver (Murray and Gridley, 2006; Tallquist and Soriano, 2000) (hereafter termed *Snai1*-cKO), the embryos still failed to develop beyond ~E9.5 (a few embryos survive until E10.5) (Lomeli et al., 2009; Murray and Gridley, 2006), indicating an essential function of this gene after gastrulation.

Analysis of E9.5 *Snai1*-cKO embryos revealed a fairly well conserved development of trunk structures. In particular, these embryos had a well-defined neural tube, midgut, notochord and a considerable number of somites that, although smaller than those of wild type littermates, retained signs of anterior/posterior compartmentalization (Figure 3 and Figure 4Ae-j). The trunk identity of this part of the *Snai1*-cKO embryos was confirmed by the presence of characteristic intermediate and lateral mesoderm tissues, including forelimb buds, molecular signatures of urogenital system development (Figure 4Ak,l) and, in some rare embryos that developed until E10.5, hindlimb buds (Figure 4Ao,p). These observations indicate that formation of the primary body is fairly well conserved in *Snai1*-cKO embryos despite *Snai1* being completely absent already at E8.0 (Murray and Gridley, 2006) when this region of the body is being laid down.

In contrast, the tail bud of E9.5 *Snai1*-cKO embryos was replaced by a protuberance protruding caudally from the trunk region of the embryo (Figure 3A,B, Figure 3—video 1, Figure 4Aa-j and Figure 4—video 1). The finding that in *Snai1*-cKO embryos *Hoxc10* expression was restricted to the protuberance is consistent with this structure replacing the tail bud (Figure 4Ac-d). A structure protruding posteriorly from the PS of *Snai1*-cKO embryos was already visible at E8.25 as a bulge (Figure 5 and Figure 5—video 1), when the embryo was building trunk structures. At this stage, *Lfng* expression in *Snai1*-cKO embryos showed variable expression patterns in the region anterior to the bulge, consistent with cycling activity in the presomitic mesoderm generating trunk somites (Figure 5Ac and Figure 5—figure supplement 1). Interestingly, variable *Lfng* signal was also observed in the bulge. To better characterize the apparently variable *Lfng* expression in the bulge we introduced the LuVeLu reporter transgene (Aulehla et al., 2008), which allows live imaging of *Lfng* cycling activity, into the *Snai1*-cKO background. Two-photon live imaging analysis of LuVeLu:: *Snai1*-cKO embryos confirmed the rather normal cyclic activity associated with trunk somite formation and revealed the existence of additional LuVeLu waves in the bulge as a signal moving posteriorly through its dorsal surface (Figure 6 and Figure 6—videos 1, 2). Therefore, *Lfng* also shows cycling activity in the bulge but with spatial and directional features different from those observed in wild type and more anterior embryonic regions of the mutant embryos, where it runs from posterior to anterior through the more ventrally located presomitic mesoderm. Together, these results indicate that the PS and the bulge represent different functional modules and that *Snai1* is required for secondary but not postcranial primary body formation.

The protuberance observed in *Snai1*-cKO embryos consisted of an epithelial-like layer extending posteriorly from the trunk neural tube, covering a mass of mesenchymal tissue contiguous with the trunk paraxial mesoderm (Figure 3, Figure 3—video 1 and Figure 5, Figure 5—video 1). At E9.5, the notochord was either bifurcated or had reversed its direction of growth (2 and 6 embryos respectively) becoming associated with gut endodermal tissue that fails to extend into the protuberance and is often detached from the rest of the embryonic tissues (found in 12 from the 19 embryos in which this feature was explored) (Figure 3Ao-r', Au-v' and B). At this stage, *Sox2* mRNA expression was detected in the trunk neural tube of *Snai1*-cKO embryos but failed to extend into the epithelial component of the bulge (Figure 3As-t). This suggests that, despite histological continuity, the bulge's epithelial sheet is most likely not an extension of the trunk neural tube. Interestingly, at E9.5 this epithelium contained *Sox2* protein [likely resulting from significantly higher protein than transcript stability (Aires et al., 2016)] in cells also expressing T (Figure 7), thus fitting the T⁺/Sox2⁺NMP molecular signature. The axial progenitor identity of the cells within this epithelium was further supported by the prominent expression of a variety of NMP-related genes (Aires et al., 2018; Rodrigo Albors et al., 2018; Gouti et al., 2017; Wymeersch et al., 2019), including *T*, *Wnt3a*, *Fgf8*, *Nkx1-2* and *Cyp26a1* (Figure 3A). Interestingly, *Fgf8* and *Nkx1-2* expression, as well as T⁺/Sox2⁺ cells, were highly restricted to the bulge's epithelial layer, which contrasts with their extension into the nascent mesenchyme in wild type embryos when they enter the tail bud stage (Figure 3A and Figure 7). These observations suggest that in the absence of *Snai1*, NMPs become trapped in the epithelial layer of the protuberance, failing to complete the trunk-to-tail transition and disrupting secondary body formation.

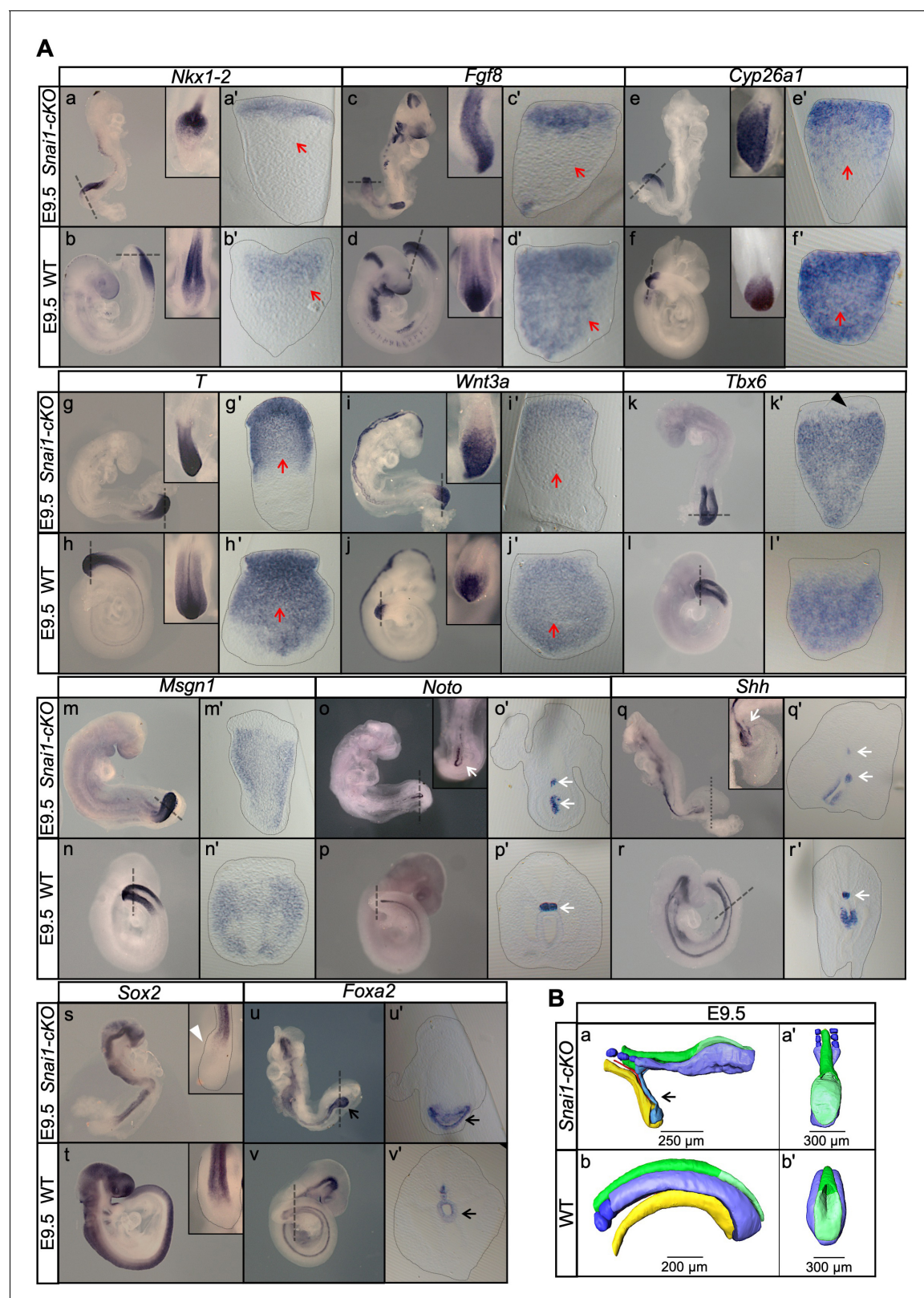


Figure 3. Axial extension is disrupted, at the level of the trunk-to-tail transition, in the absence of *Snai1*. **(A)** Wholemount in situ hybridization with the indicated probes in E9.5 wild type (WT) and *Snai1*-cKO embryos. Axial progenitor-related markers *Fgf8*, *Nkx1-2*, *Cyp26a1* were abnormally restricted to the epithelium of the caudal protuberance of *Snai1*-cKO embryos (red arrows) and *T* and *Wnt3a*-stained embryos were substantially downregulated in the central component of the bulge mesenchyme. *Tbx6* expression was observed in the epithelial-like component of the bulge (black arrowhead) in Figure 3 continued on next page

Figure 3 continued

addition to its mesenchymal expression. *Msgn1* was present in the presomitic mesoderm and in the mesenchymal component of the bulge. The white arrows in the *Noto*-stained embryos indicate the bifurcated or inverted notochord growth in the absence of *Snai1*. *Sox2* was absent from the bulge region (white arrowhead) of *Snai1*-cKO embryos. Black arrows in the *Foxa2* labelled embryos highlight the abnormal localization of hindgut endoderm in *Snai1*-cKO embryos. (B) 3D reconstructions of E9.5 *Snai1*-cKO and WT posterior/caudal structures: neural tube (green), open epiblast (light green), presomitic mesoderm (light blue), somites (dark blue), notochord (red) and endoderm (yellow). At this stage, the ectopic bulge of *Snai1*-cKO embryos forms a structure that resembles an abnormally extended open epiblast in which the closing neural plate fails to extend caudally. The notochord often bifurcates, with one end following the posterior gut endoderm that is detached from the rest of the embryonic structures (black arrow).

The online version of this article includes the following video for figure 3:

Figure 3—video 1. Animated 3D reconstruction of the tail bud of E9.5 embryos.

<https://elifesciences.org/articles/56615#fig3video1>

The mesenchymal component of the bulge was positive for paraxial mesoderm markers, including *Tbx6* or *Msgn1*, but negative for lateral mesoderm markers such as *Tbx4* (Figure 3Ak-n', Figure 4A and Figure 5Ac-d'), thus resembling paraxial mesodermal features. Moreover, *Tbx6* expression was not restricted to the bulge's mesenchyme but was also observed in the epithelium (Figure 3Ak-l', and Figure 8B). Since a portion of tail bud NMPs are positive for *Tbx6* expression (Javali et al., 2017), it is possible that the bulge's epithelial component contains progenitor cells that have already acquired some of the mesenchymal traits associated with tail bud axial progenitors. Consistent with this hypothesis, we only detected residual *Cdh1* and *Epcam* expression in the bulge epithelium at E9.5, whereas expression of mesenchymal markers such as *Cdh2* and *Vim* were readily detectable at levels comparable to those in wild type embryos (Figure 8A and Figure 8—figure supplement 1). In addition, *Laminin1* expression in the bulge's epithelium was highly disorganized already at E8.5, contrasting with the characteristic epithelial pattern observed in adjacent more anterior areas of the same embryos or in the epiblast of wild type littermates (Figure 8B). Together, these observations indicate that in the absence of *Snai1* axial progenitors initiate tb-EMT and acquire some mesenchymal features but are unable to complete the transition to tail bud development.

***Snai1* and *Tgfb1* cooperatively orchestrate the transition from primary to secondary body formation**

Gdf11 has been associated with the trunk-to-tail transition (Aires et al., 2016; Jurberg et al., 2013; Liu, 2006; Matsubara et al., 2017). However, its partial functional redundancy with *Gdf8* in this process (McPherron et al., 2009; McPherron et al., 1999) hinders proper evaluation of its contribution to tb-EMT. Nevertheless, genetic experiments indicate that *Gdf11* (and most likely *Gdf8*) activity in the caudal embryo is mediated by *Tgfb1* (also known as *Alk5*) (Andersson et al., 2006; Jurberg et al., 2013), a known EMT regulator (Derynck et al., 2014; Xu et al., 2009) that is expressed in areas containing axial progenitors (Figure 2—figure supplement 2), thus making *Gdf11/Tgfb1* signalling a prime candidate to play a role in tb-EMT. To overcome *Gdf11/Gdf8* functional redundancy, we therefore generated *Tgfb1* mutant embryos. Initial analysis of *Tgfb1*^{-/-} embryos confirmed the role of this receptor as an inducer of the trunk-to-tail transition (Figure 9), and also indicating the absence of redundancy from other type I receptor of the *Tgfb/BMP* gene family in this process. Indeed, these mutant embryos extended their axis through the trunk region but failed to undergo the transition to tail development, as illustrated by absent hind limb bud markers, lack of *Lin28a* expression at E10.5, which is involved in promoting tail bud axial progenitor activity (Aires et al., 2019; Robinton et al., 2019), and only residual activation of posterior genes like *Hoxc10* (Figure 9). In addition, the anterior-ventral relocation of the allantois -as the embryo turns and starts developing umbilical structures at tail bud stages- was compromised in these mutant embryos and all tested NMP markers (e.g. *T*, *Fgf8* and *Cyp26a1*) seemed to stay associated with dorsal tissues, with no sign of contribution to tail bud mesenchyme/mesodermal tissue (Figure 9G-L). Together, these observations are consistent with the embryos finishing their axial extension when *Pou5f1* stops influencing axial progenitor activity derived from its natural decay (Osorno et al., 2012), as they are unable to activate the trunk-to-tail transition.

In *Tgfb1*^{-/-} embryos the transition from *Cdh1* to *Cdh2* in the posterior embryo was very inefficient, as *Cdh1* was still detected in the *Tgfb1*^{-/-} epiblast at E9.5 and *Cdh2* was only detected at very low levels (Figure 10A). Concomitantly, we observed the characteristic *Laminin1* epithelial patterns,

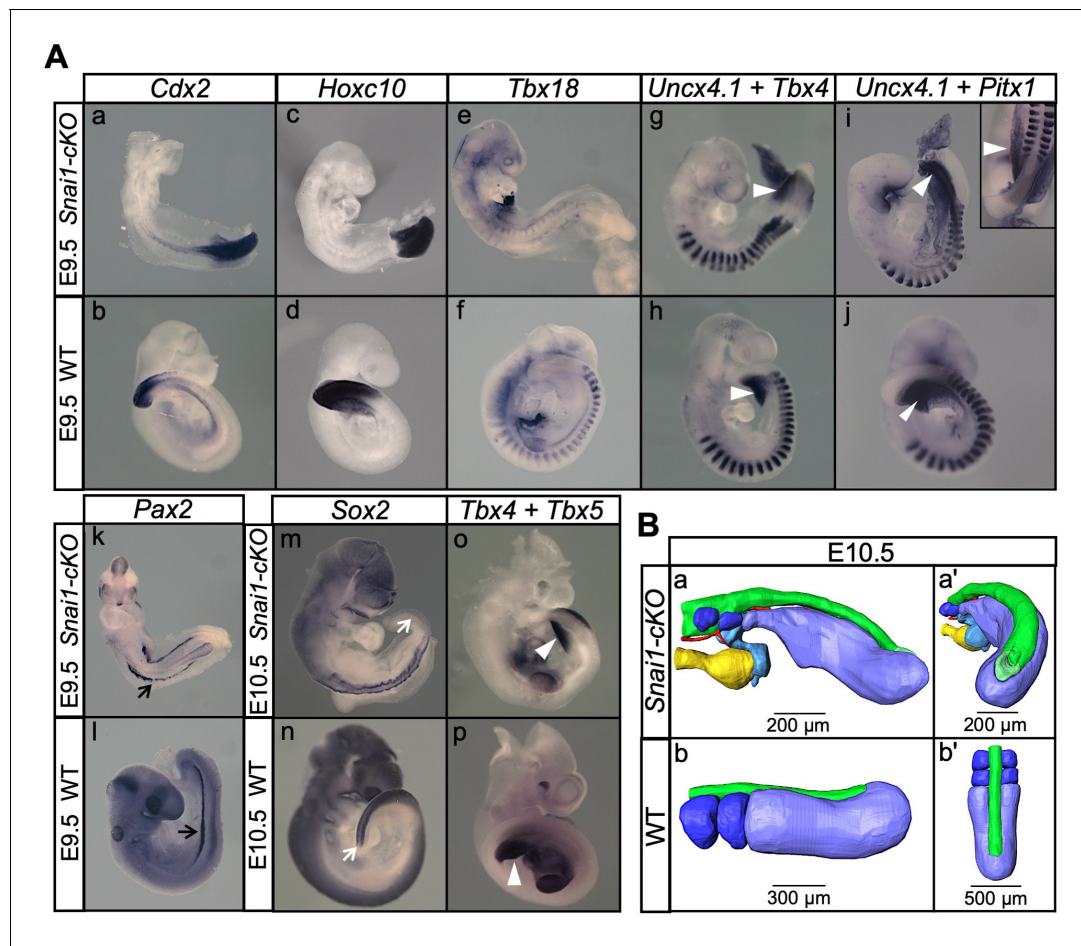


Figure 4. *Snai1*-cKO embryos develop fairly-well organized primary body structures but are unable to form a tail bud. (A) Wholemount in situ hybridization with the indicated markers in wild type (WT) and *Snai1*-cKO mutant embryos at E9.5 and E10.5. The insets show close ups of the caudal region. Transverse sections at the levels indicated in the whole-mount embryo are also shown. *Cdx2* and *Hoxc10* expression in *Snai1*-cKO embryos indicate that the bulge is the equivalent of the tail bud in similar stage wild type embryos. The black arrows in the *Pax2*-stained embryos indicate the presence of intermediate mesoderm in both *Snai1*-cKO and wild type embryos. At E9.5, *Tbx18* expression and the combined in situ hybridization for *Uncx4.1/Tbx4* and *Uncx4.1/Pitx1*, indicate the presence of fairly-well organized trunk somites and hindlimb buds (white arrowheads) in *Snai1*-cKO embryos, which is also highlighted in some mutant embryos, that survived at E10.5, with *Tbx5* and *Tbx4* expression. White arrows indicate that *Sox2* is still downregulated in the bulge of E10.5 *Snai1*-cKO embryos, in comparison to the tail bud of wild type littermates. (B) 3D reconstructions of caudal structures of E10.5 wild type and *Snai1*-cKO embryos. Wild type embryos have a closed neural tube (in green), dorsal to the presomitic mesoderm (in blue) and somites (in dark blue). In contrast, *Snai1*-cKO embryos have a bifurcated notochord (red), detached endoderm (yellow) and the bulge resembles a still open caudal epiblast (in green). Somites are shown in dark blue and extra mesoderm in *Snai1*-cKO embryos is highlighted in cyan. Note that the structures posterior to the last somites in *Snai1*-cKO embryos did not extend further than at E9.5, leading to a shorter axis.

The online version of this article includes the following video for figure 4:

Figure 4—video 1. Animated 3D reconstruction of the tail bud of E10.5 embryos.

<https://elifesciences.org/articles/56615#fig4video1>

as well as higher *Epcam* and lower *Vim* levels in the *Tgfbf1* mutant epiblast than in similar stage control embryos (**Figure 10B** and **Figure 10—figure supplement 1**). These data support an essential role for *Gdf11/Tgfbf1* signalling in triggering tb-EMT and suggest that this signalling activity precedes that of *Snai1* during tb-EMT. However, contrary to what has been described in other experimental contexts (**Lamouille et al., 2014**), *Snai1* is most likely not a direct target of *Tgfbf1* signalling during tb-EMT, as we did not find any evidence for increased *Snai1* expression in the progenitor-containing region of *Tgfbf1*-overexpressing embryos [*Cdx2-Alk5^{CA}* transgenics (**Jurberg et al., 2013**; **Figure 11B**)]. This suggests that *Snai1* and *Gdf11/Tgfbf1* signalling are part of independent pathways that converge to orchestrate the tb-EMT. Indeed, *Snai1* expression in the PS region covers

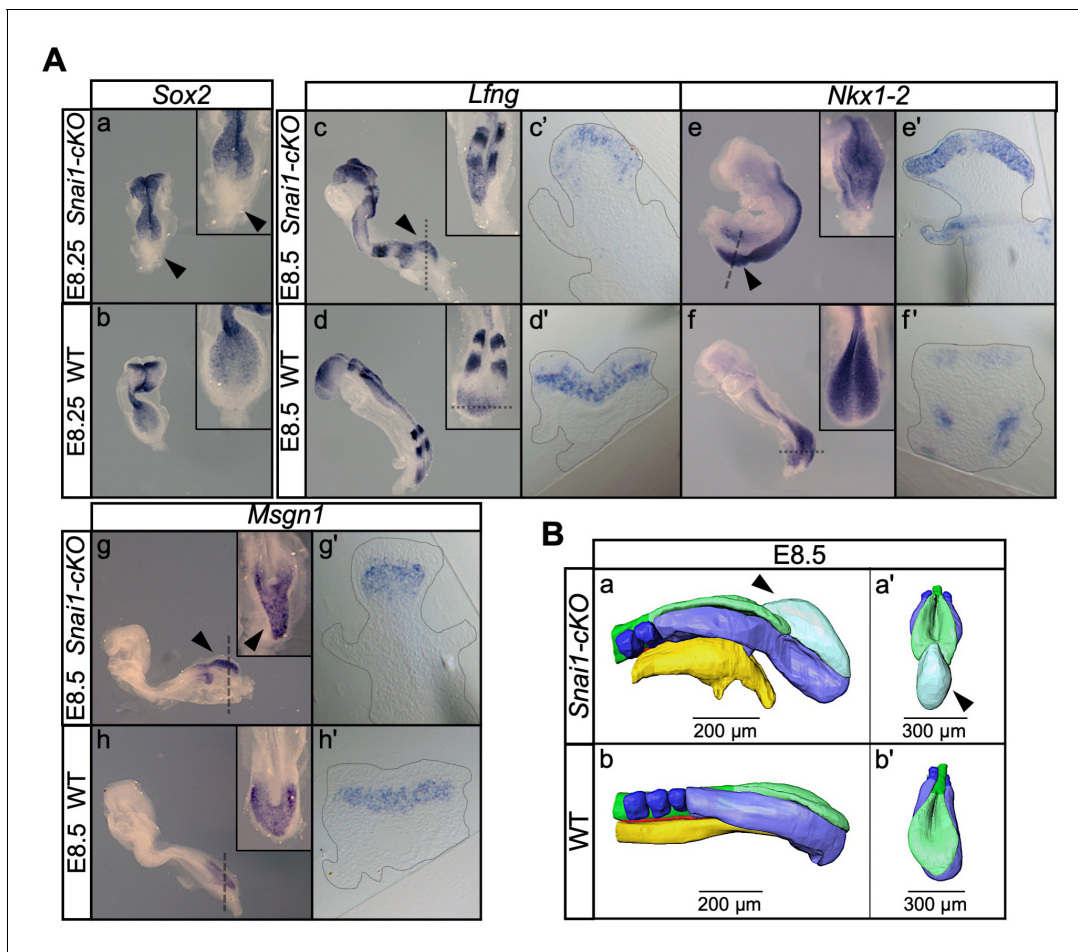


Figure 5. *Snai1*-cKO embryos develop an ectopic bulge associated with the PS. (A) Wholemount in situ hybridization with the indicated markers in wild type (WT) and *Snai1*-cKO mutant embryos at E8.5. The insets show close ups of the caudal region. Transverse sections at the levels indicated in the whole-mount embryo are also shown. In the absence of *Snai1*, a protuberance (black arrowheads) starts to arise from the PS around E8.25. This ectopic bulge is positive for *Lfng* and *Nkx1-2*, but not for *Sox2*. *Lfng* and *Nkx1-2* expression are restricted to the epithelial-part of the bulge. *Msgn1* was only found in the mesenchymal component. (B) 3D reconstruction of E8.5 *Snai1*-cKO and wild type posterior/caudal structures: neural tube (green), open epiblast (light green), presomitic mesoderm (light blue), somites (dark blue), notochord (red) and endoderm (yellow). At E8.5, *Snai1*-cKO embryos contain an ectopic bulge (black arrowhead) associated with the PS and the hindgut endoderm shows abnormal development.

The online version of this article includes the following video and figure supplement(s) for figure 5:

Figure supplement 1. *Lfng* dynamic activity in the bulge of *Snai1*-cKO embryos.

Figure 5—video 1. Animated 3D reconstruction of the tail bud of E8.5 embryos.

<https://elifesciences.org/articles/56615#fig5video1>

only a small region within the *Gdf11* expression domain in the epiblast (Hernández-Martínez et al., 2019; Nieto et al., 1992; Figure 11A). Therefore, it is possible that *Snai1* marks a subset of axial progenitors within the cell pool exposed to *Gdf11*/*Tgfb β 1* signalling to be recruited to the tail bud by entering tb-EMT.

Extended *Snai1* expression in the PS mobilizes functional tail bud progenitors

A prediction from this hypothesis is that expanding the *Snai1* expression domain in the PS would lead to an increase in the number of cells entering the tail bud route. We tested this hypothesis using a transgenic approach expressing *Snai1* under the control of the *T-str* enhancer (Clements et al., 1996), a regulatory element of the *Brachyury* gene that drives expression in progenitor cells within the PS (Anderson et al., 2013). About half of these transgenic embryos (n = 43)

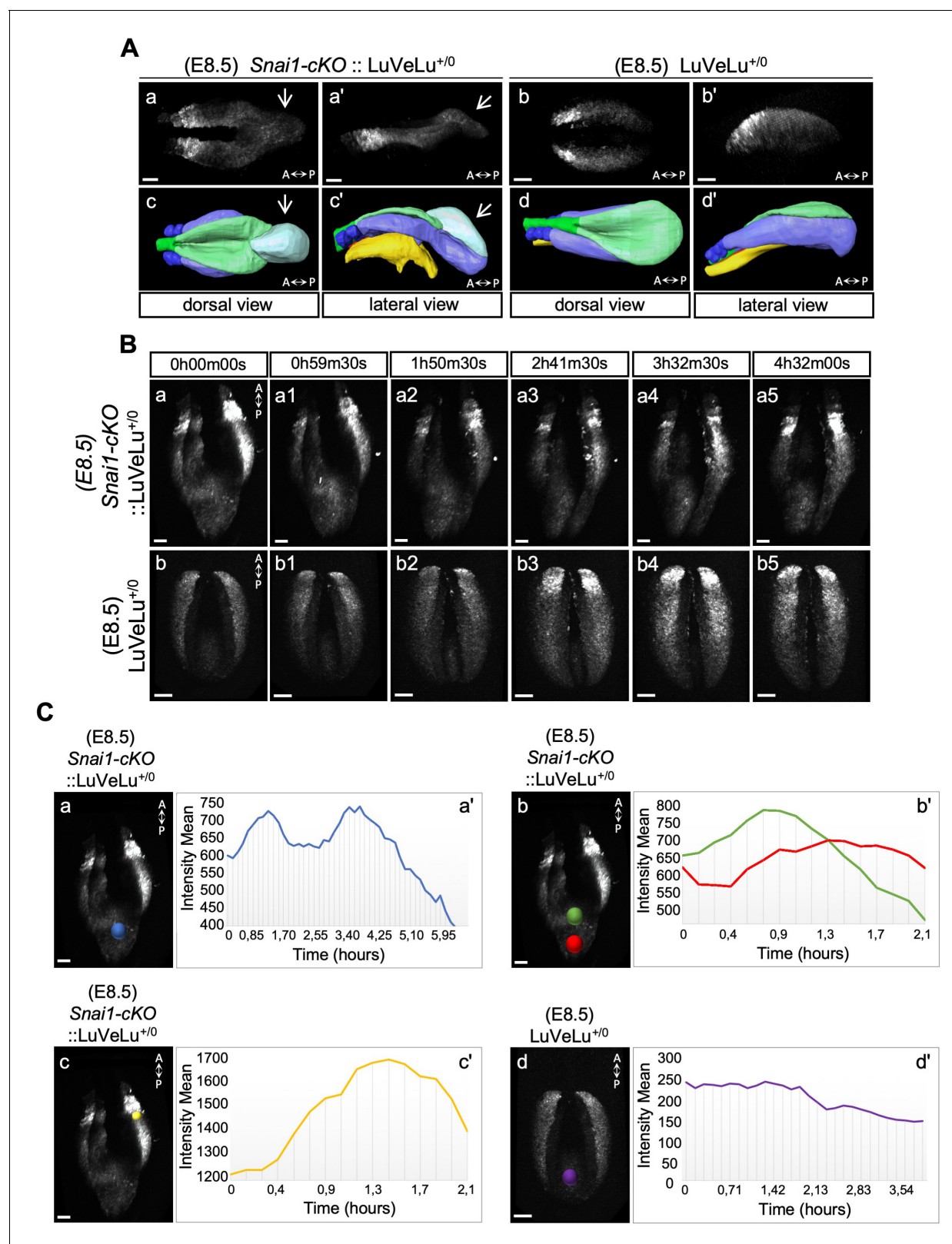


Figure 6. 3D two-photon live imaging of LuVeLu reporter expression in E8.5 *Snai1-cKO* and control embryos. (A) Snapshot at time-point = 0 of the LuVeLu reporter, in *Snai1-cKO* (from **Figure 6—video 1**) and control (from **Figure 6—video 2**) E8.5 embryos. In addition to the normal LuVeLu signal in the presomitic mesoderm, *Snai1-cKO* embryos display ectopic LuVeLu expression in the epithelial-like component of the bulge. (B) Snapshots from **Figure 6—video 1** and **Figure 6—video 2**, of time-lapsed two-photon live imaging of the LuVeLu reporter in E8.5 *Snai1-cKO* and control embryos at

Figure 6 continued on next page

Figure 6 continued

the indicated time points. *Snai1*-cKO embryos have ectopic LuVeLu expression in the epithelial-like component of the bulge in addition to the normal LuVeLu signal in the presomitic mesoderm. (C) Quantitative analysis of LuVeLu cycling activity in the bulge of *Snai1*-cKO embryos. Intensity mean was calculated in the region highlighted by the blue spot (Ca) and plotted for each imaged time-point (8.5 min interval; Ca'). The existence of two-peaks (at $t = 1.3$ hr and $t = 3.6$ hr of the time-lapse) and a substantial decrease between them, suggests cycling activity in the bulge of the mutant embryo. These waves occur from anterior to posterior, once the higher intensity mean peak observed in the posterior part of the bulge (red spot; Cb) occurred later than the higher peak measured in the anterior part of the bulge (green spot; Cb). The higher measured wave peak in the posterior part of the bulge (red spot; Cb) coincides with the time-point corresponding to the higher intensity mean peak detected in the newly formed somite (yellow spot; Cc). No signs of LuVeLu cycling activity were observed near the PS (purple spot; Cd) in LuVeLu^{+/0} control embryos. Scale bar: 50 μ m.

The online version of this article includes the following video(s) for figure 6:

Figure 6—video 1. LuVeLu activity in a E8.5 *Snai1*-cKO embryo.

<https://elifesciences.org/articles/56615#fig6video1>

Figure 6—video 2. LuVeLu activity in a E8.5 wild type embryo.

<https://elifesciences.org/articles/56615#fig6video2>

showed caudal morphological abnormalities at E9.5, including axial shortening and premature closure of the caudal neuropore (**Figure 12**). In these embryos the *T*, *Cyp26a* and *Nkx1.2* domains expanded into the ventral mesenchymal region of the shortened tail bud (**Figure 12C-D2, I-L2**), consistent with production and delamination of tail bud axial progenitors in regions more anterior to the normal position of the tail bud. In addition, these embryos had a variable number of ectopic neural tubes connected caudally with the expanded *Nkx1.2*-positive mesenchymal domain, aligning ventrally to the main spinal cord (**Figure 12**), which is consistent with these cells having acquired tail bud NMP identity. We also found cells expressing *T* and *Tbx6* intermingled with these ectopic neural tubes (**Figure 12C1-C2, M1**), further supporting the NMP identity of these progenitors. Interestingly, these extra neural tubes contained a degree of dorso-ventral patterning, as shown by *Foxa2* expression in their ventral domain, corresponding to the floor plate (**Figure 12O1**), and by *Wnt3a* expression in the dorsal part of some of these ectopic neural tubes (**Figure 12G1**). Together, these results indicate that *Snai1* is necessary and sufficient for axial progenitor mobilization during tail bud formation.

Discussion

The transition from primary to secondary body formation involves at least two coordinated but independent processes. One such process entails activation of a terminal differentiation program in the progenitors for the intermediate and lateral components of the mesoderm that results in the induction of the hindlimb bud and of the mesodermal component of the cloaca (*Jurberg et al., 2013*). The second process involves the common progenitors for the spinal cord and paraxial mesoderm that become the tail bud, from where they promote further extension of the body axis to generate post-sacral body structures (*Henrique et al., 2015; Steventon and Martinez Arias, 2017; Wilson et al., 2009*).

Snai1 and *Tgfb1* play a key role in the gene regulatory networks driving tail bud formation

Genetic experiments have shown that *Gdf11* plays an important role in both processes by activating signalling in the axial progenitors through the *Tgfb1* receptor. Indeed, premature activation of this signalling in axial progenitors anticipates the trunk-to-tail transition (*Jurberg et al., 2013; Liu, 2006; Matsubara et al., 2017*), and *Tgfb1* null embryos are unable to engage in tail development (this work).

A variety of experimental evidence indicates that *Isl1* is a key component of the network downstream of *Tgfb1* to induce hindlimb and cloacal tissues (*Jurberg et al., 2013*). The same experiments also showed that this gene is not involved in tail bud formation, indicating the existence of a different mechanism regulating the formation of this structure. Here we presented evidence for the critical role of *Snai1* in this process. Interestingly, however, this gene is not under the control of *Tgfb1* signalling but provides instead a parallel activity that promotes segregation of axial progenitors into the future tail bud upon functional convergence with *Tgfb1* signalling. The *Snai1*-cKO

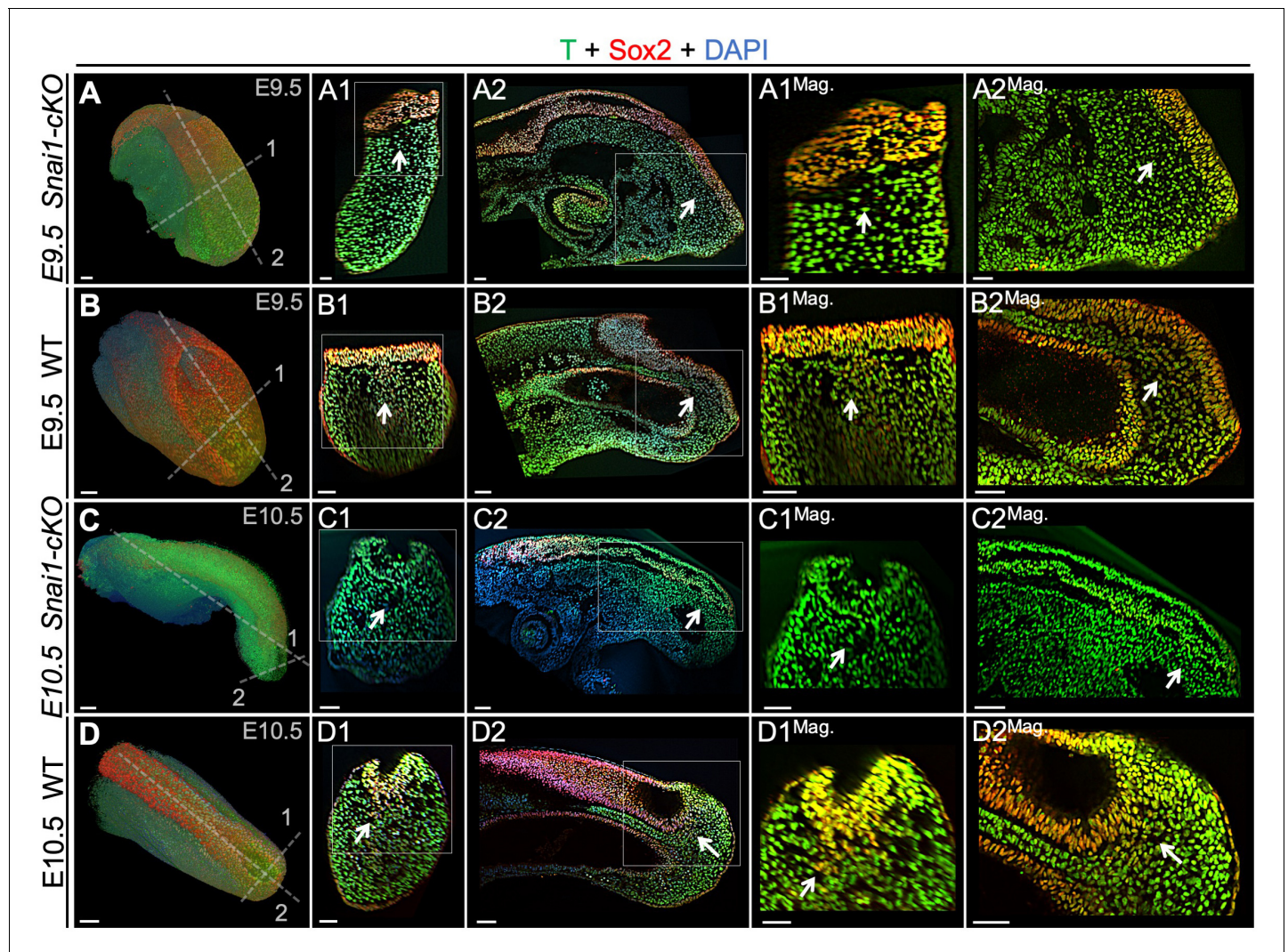


Figure 7. T and Sox2 double-positive cells are restricted to the epiblast of *Snai1-cKO* embryos. Wholemount immunostaining for T (green), Sox2 (red) and DAPI (blue) in E9.5 (A and B) series and E10.5 (C and D) series *Snai1-cKO* and wild type (WT) embryos. Transversal and sagittal optical sections together with magnifications (Mag.) are also shown. In E9.5 wild type embryos, T⁺/Sox2⁺ cells are found in the mesenchyme below the epiblast (white arrows). Conversely, in similar staged *Snai1-cKO* embryos, they are restricted to the epithelial component of the bulge. At E10.5 most of the bulge's epithelium, of *Snai1-cKO* embryos, closed into a tube that is mostly negative for Sox2 and positive for T. In contrast, T and Sox2 double-positive cells are still found in mesenchymal compartments of the tail bud of wild type embryos. Magnifications are shown without DAPI. Scale bars correspond to 50 μ m.

mutant phenotype indicates that tail bud formation starts rather early in development when the embryo is still actively engaged in PS-mediated trunk development, much earlier than the appearance of morphological signs of a tail bud. This phenotype thus uncovers the existence of a functional developmental module specifically devoted to the formation of the tail bud, independent from that involved in trunk formation. The existence of such 'tail bud module' somehow resembles hindlimb or cloacal induction, thus suggesting a unifying mode by which Gdf11/Tgfb β 1 signalling activity triggers the different processes involved in the trunk-to-tail transition, consisting in activating alternative developmental programs in different subsets of axial progenitors. The observation that NMPs are suppressed by *Isl1* activity (Jurberg *et al.*, 2013) indicates that the intrinsic properties of these cells differ from those of the progenitors for intermediate and lateral mesoderm, determining whether they engage in tail bud or hindlimb/cloaca formation upon Tgfb β 1 functional input.

Different experimental observations indicate that trunk length and the position of the switch from trunk to tail developmental modes result from quantitative functional balance between Gdf11 and

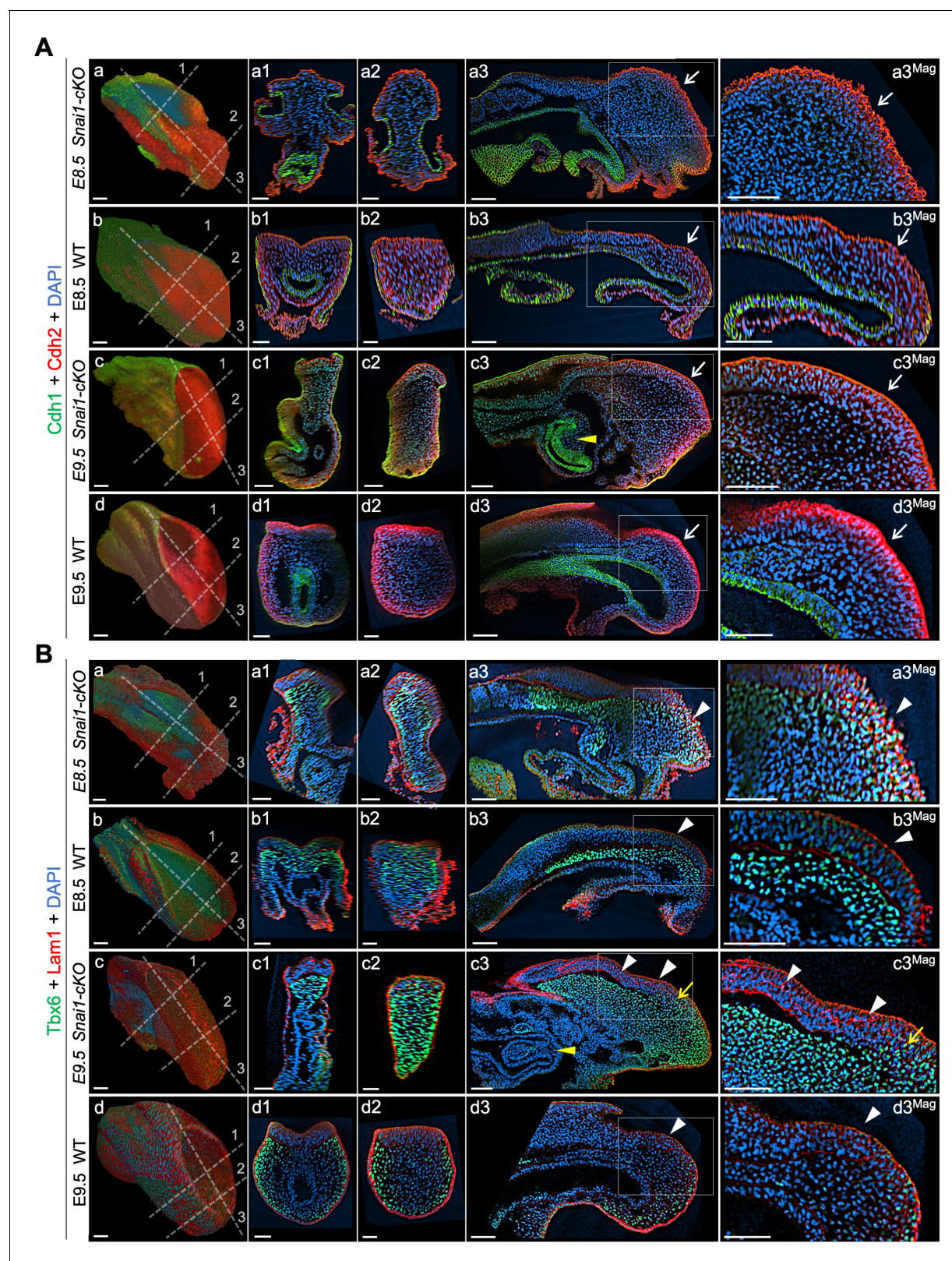


Figure 8. Axial progenitors acquire mesenchymal properties in the absence of *Snai1*. Wholemount immunostainings for *Cdh1* (green) plus *Cdh2* (red) (A) and *Tbx6* (green) plus *Lam1* (red) (B), in E8.5 and E9.5 *Snai1*-cKO and wild type (WT) embryos. Transversal and sagittal optical sections in the indicated regions are also shown together with magnifications (Mag.). The transition from *Cdh1* to *Cdh2* still occurs in the epithelial component of the bulge in the absence of *Snai1* (white arrows). This region of *Snai1*-cKO embryos contains ectopic *Tbx6*-positive cells (yellow arrow) and a severely

Figure 8 continued on next page

Figure 8 continued

disorganized *Lam1* expression (white arrowheads). Yellow arrowheads highlight posterior gut bifurcation in E9.5 *Snai1*-cKO embryos. Scale bars: 50 μ m. DAPI staining in blue.

The online version of this article includes the following figure supplement(s) for figure 8:

Figure supplement 1. Wholemount immunohistochemistry in wild type and *Snai1*-cKO embryos.

Pou5f1 activities (Aires et al., 2016; DeVeale et al., 2013; Jurberg et al., 2013; Mallo, 2018). These observations, together with the *Snai1*-cKO mutant phenotype lead us to suggest a model for axial extension whereby at early stages (mouse E8.0) progenitor activity is dominated by *Pou5f1*, keeping an active PS, while *Gdf11*/*Tgfb β 1* functional contribution is negligible or extremely small. As

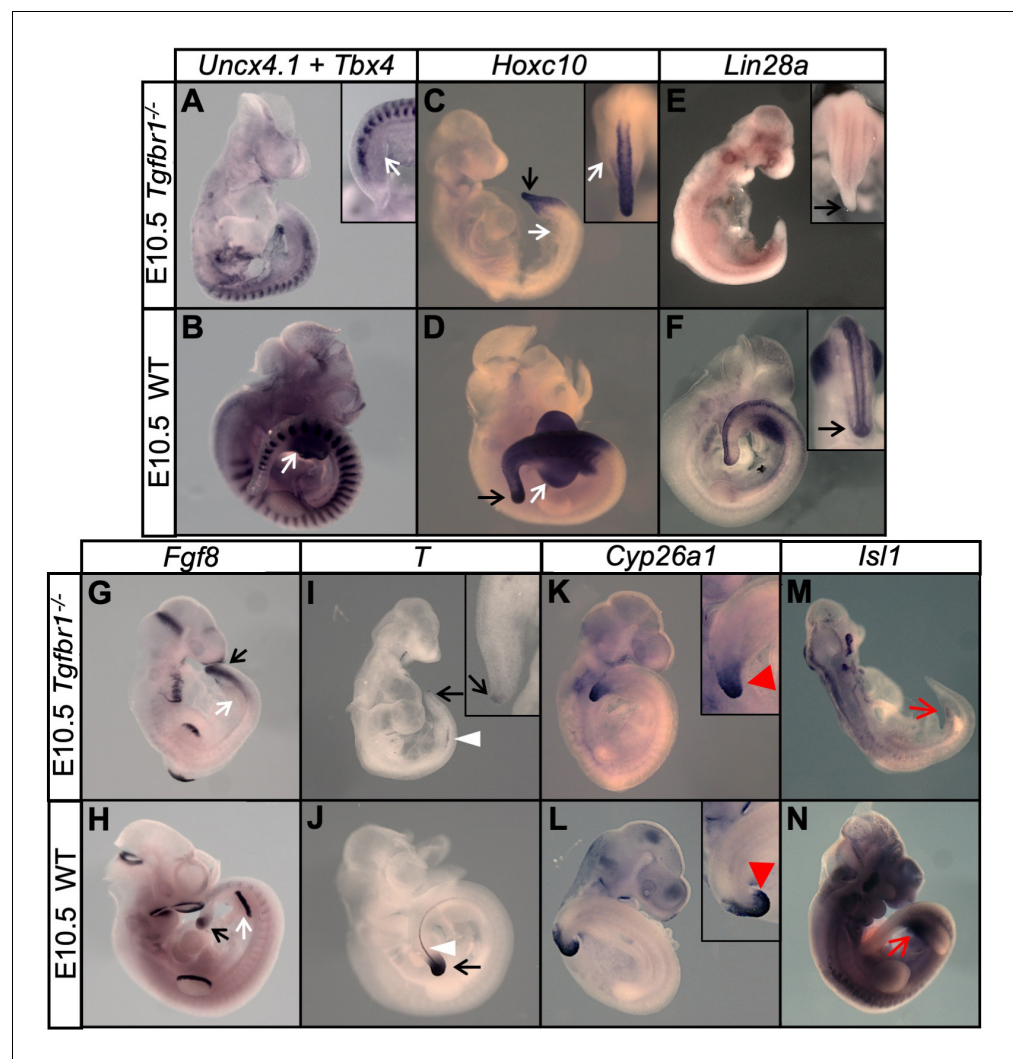


Figure 9. *Tgfb β 1* mutant embryos are unable to undergo the trunk-to-tail transition. (A–N) Wholemount in situ hybridization with the indicated markers in E10.5 *Tgfb β 1*^{-/-} and wild type (WT) embryos. Absent hindlimb bud formation in *Tgfb β 1* mutants is highlighted by absent *Tbx4*, *Fgf8* and *Hoxc10* expression (A–D, G and H) in the relevant area (white arrows). Tail bud formation is deficient in *Tgfb β 1* mutant embryos (black arrows). *Lin28a* and *T* are drastically downregulated in the tail of *Tgfb β 1* mutant embryos (black arrows in E and I), suggesting that these mutants are caudally truncated at this stage. The white arrowheads indicate the notochord, which did not continue its development to more caudal regions in *Tgfb β 1* mutant embryos. Red arrowheads highlight the abnormal *Cyp26a1* expression in the *Tgfb β 1*^{-/-} tail. Red arrows in (M) and (N) indicate absent *Isl1* expression in the prospective developing cloaca of *Tgfb β 1* mutant embryos.

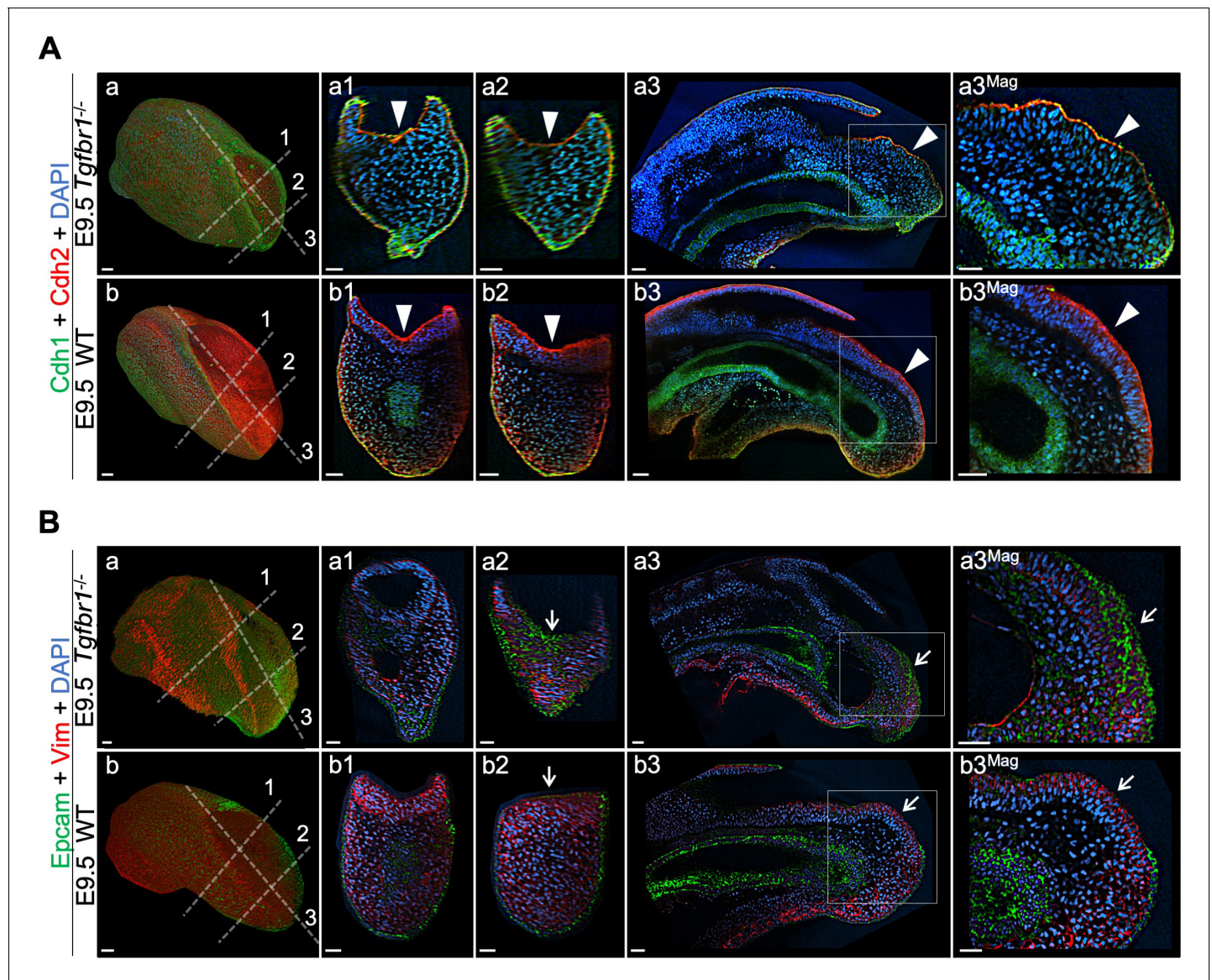


Figure 10. Tail bud EMT is compromised in the absence of *Tgfb1*. Wholemount immunostaining for Cdh1 (green) plus Cdh2 (red) (A) and Epcam (green) plus Vim (red) (B) in *Tgfb1*^{-/-} and wild type (WT) embryos. Transversal and sagittal optical sections through the indicated regions are also shown together with magnifications (Mag.). White arrowheads indicate incomplete Cdh1 to Cdh2 switch in *Tgfb1* mutants. White arrows show persistent Epcam and deficient Vim expression in the epithelium of the mutant embryos. DAPI staining is shown in blue. Scale bars: 50 μm.

The online version of this article includes the following figure supplement(s) for figure 10:

Figure supplement 1. Whole mount immunohistochemistry in wild type and *Tgfb1* embryos.

development proceeds *Pou5f1* activity is progressively reduced in the epiblast (Osorno et al., 2012) and *Tgfb1* acquires increased functional weight in this area. *Snai1*-positive cells in the PS are the first to respond to the reversed *Pou5f1*/*Tgfb1* activity balance, resulting in NMPs mobilizing to the tail bud. This process is already triggered when the PS still actively coordinates development at more anterior levels, indicated by the *Snai1*-cKO phenotype. As development proceeds, *Pou5f1* activity decreases (Osorno et al., 2012), and given its requirement for PS maintenance (Aires et al., 2016; DeVeale et al., 2013), it is likely to be responsible, together with *Snai1*, for PS regression and caudal epiblast closure. Under these conditions, *Tgfb1* eventually takes full control of axial progenitor activity, leaving the tail bud as the only source of axial extension activity and recruiting lateral mesoderm progenitors into hindlimb and cloacal structures. In addition, this model implies an

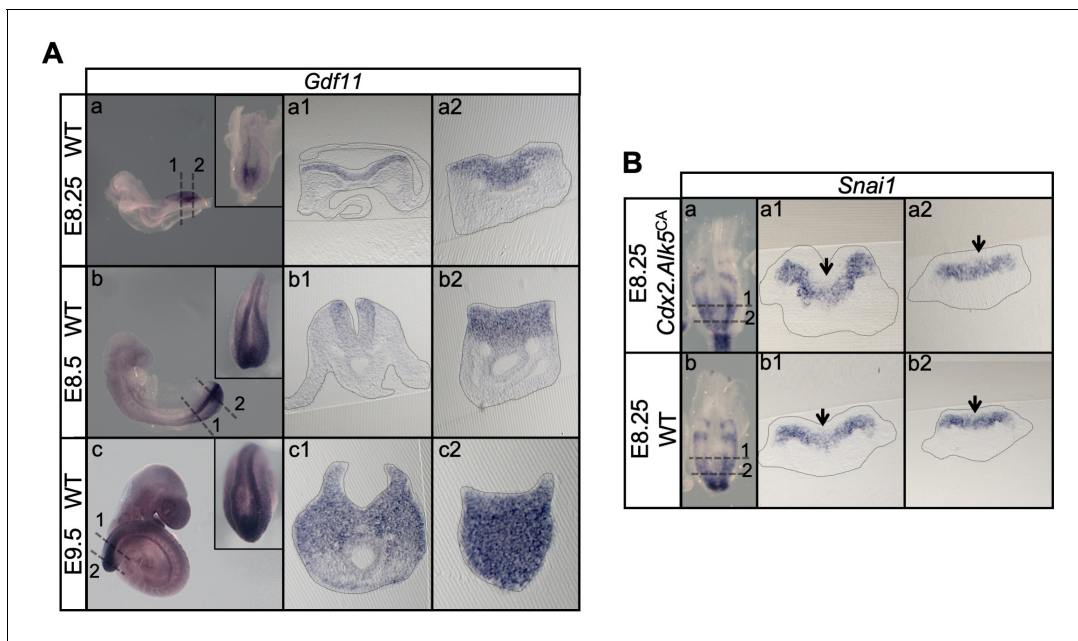


Figure 11. *Snai1* is not a downstream target of *Tgfb β 1*. (A) Wholemount in situ hybridization for *Gdf11* in E8.25, E8.5 and E9.5 wild type (WT) embryos. Transversal sections through the areas indicated in the whole mounted embryo highlight the increase of *Gdf11* in caudal tissues during primary body formation. (B) Overexpression of a constitutively active form of *Tgfb β 1* (transgenic *Cdx2-Alk5^{CA}*) does not result in increased *Snai1* expression as shown in transversal sections of the areas indicated in the wholemount embryo. Arrows indicate the position of the PS, where low levels of *Snai1* expression can be observed both in the WT (Bb) and transgenic embryos (Ba).

overlap of primary and secondary processes of neural tube formation, and therefore fits better with classical observations in human embryos (Saitsu et al., 2004). A failure in the coordination between these two processes can ultimately lead to the generation of spina bifida (Saitsu and Shiota, 2008).

Two different functional types of EMT are required during vertebrate axial extension

Here we have shown that the transition from primary to secondary body formation in the mouse entails an incomplete EMT acting on axial progenitors and termed this new developmental process 'tail bud EMT' (tb-EMT), as it is required for tail bud formation. This EMT is different from that driving epiblast progenitors through the PS to generate mesodermal tissues during gastrulation in different ways. (1) progenitors undergoing tb-EMT do not enter a differentiation route but retain instead progenitor properties that endow them with the capacity to further extend the body axis by generating both the neural tube and paraxial mesoderm of the tail. (2) tb-EMT is incomplete, keeping expression of a subset of epithelial markers that leave the progenitors in a transitional state. This property might actually facilitate the production of both neural and mesodermal structures from tail bud axial progenitors. Moreover, it is possible that maintenance of some epithelial properties can help progenitors to keep contact with the trunk neural epithelium that they will eventually extend by regaining full epithelial features. Conversely, these cells would be able to enter mesodermal differentiation routes just by completing the EMT, thus circumventing the need for a functional PS. Interestingly, comparison of CNH and tail bud mesenchyme heterochronic grafts (Cambray and Wilson, 2002) showed that after tail bud progenitors enter the mesodermal compartment, they have no longer the potential to generate neural tissues, becoming restricted to mesodermal compartments, further supporting that tail bud progenitors are kept in a transitory state instead of acquiring full mesenchymal characteristics when entering the tail bud.

Another major difference between gastrulation- and tail bud- associated EMTs resides in their genetic control. Genetic experiments identified a number of genes involved in early gastrulation steps, including *Crumbs2*, *p120-catenin*, *Eomes*, *Nodal* and FGF signalling (Arnold et al., 2008; Deng et al., 1994; Hernández-Martínez et al., 2019; Ramkumar et al., 2016; Voiculescu et al.,

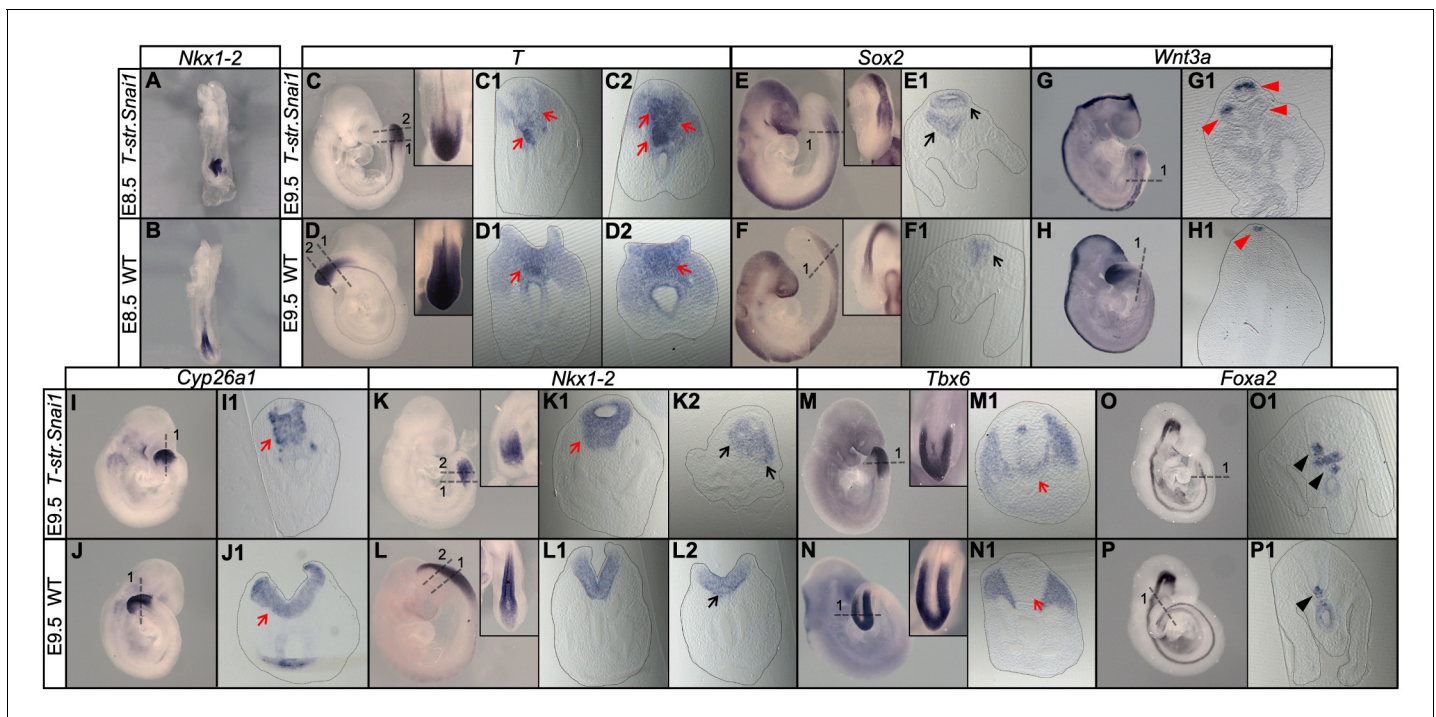


Figure 12. Extended *Snai1* expression in the PS is sufficient to mobilize axial progenitors from the epiblast. (A–R) Wholemount in situ hybridization for the indicated markers in wild type (WT) and transgenic *T-str-Snai1* embryos. *Nkx1-2* expression in the transgenic embryos (A) indicates an already deformed caudal epiblast at E8.5. At E9.5 *T-str-Snai1* embryos are caudally truncated and display a complete premature closure of their caudal epiblast. Red arrows in C2, I1 and K1 highlight the ventrally extended *T*, *Cyp26a1* and *Nkx1-2* expression, indicating ectopic mobilization of axial progenitors. Black arrows in E1 and K2 show ventrally located ectopic neural tubes in *T-str-Snai1* embryos. Red arrows in C1 and M1 indicate *T*, *Tbx6*-positive cells intermingled with the ectopic neural tubes. The black arrowheads in O1 and P1 indicate the floor plate, showing conserved ventral patterning in ectopic neural tubes of *T-str-Snai1* embryos. The red arrowheads in G1 show *Wnt3a* expression in the dorsal region of the ectopic neural tubes.

2014; Yamaguchi et al., 1994). *Tgfr1*, which is an essential component of tb-EMT, is clearly not involved in gastrulation as embryos lacking this receptor can go through this developmental step and remarkably generate trunk structures. *Snai1* seems to be required for both gastrulation and tb-EMT, although its role might be different in both processes. *Snai1* null mutant embryos generate mesoderm during gastrulation but the mesodermal cells produced are unable to complete their EMT process, keeping several epithelial characteristics and failing to downregulate *Cdh1* (Carver et al., 2001). During tb-EMT cells lacking *Snai1* had already lost several epithelial characteristics, most likely resulting from a previous *Tgfr1* functional input, suggesting that its role during tail bud formation must be different from that in gastrulation. In *Snai1*-cKO embryos, axial progenitors were restricted to epithelial-like layer of the bulge contiguous with the posterior epiblast. Conversely, these cells seem to leave the epiblast prematurely in *T-str-Snai1* transgenic embryos. Therefore, although the role of *Snai1* in tail bud formation requires further evaluation, it is possible that it might be involved in cell mobilization. Interestingly, the phenotypes of both *Snai1* null and of *Snai1*-cKO mutant embryos suggest that PS activity during postcranial elongation is functionally different to that during early gastrulation because *Snai1*-cKO embryos are able to efficiently elongate the postcranial primary body even in the absence of detectable *Snai1* (Murray and Gridley, 2006). This observation is in keeping with a large body of genetic data indicating differences in the gene networks regulating early gastrulation processes and postcranial axial elongation (Andre et al., 2015; Herrmann et al., 1990; Liu et al., 1999; Savory et al., 2011; Takada et al., 1994).

It should be also noted that tb-EMT functional and molecular characteristics are more akin to those described for the transitions involved in metastatic processes than to the typical developmental EMTs (e.g. gastrulation) (Acloque et al., 2009; Kalluri and Weinberg, 2009; Lamouille et al., 2014; Nieto et al., 2016). These transient tail bud progenitors might not only be used to re-assess

neural tube closure defects, but also represent a novel in vivo model to study the mechanisms activating *Tgfr1*/*Snai1*-dependent metastatic processes.

Materials and methods

Key resources table

Reagent type (species) or resource	Designation	Source or reference	Identifiers	Additional information
Gene (<i>M. musculus</i>)	<i>Snai1</i>	MGI	MGI:98330	(other names) <i>Snail</i>
Gene (<i>M. musculus</i>)	<i>Tgfr1</i>	MGI	MGI:98728	(other names) <i>Alk5</i>
Genetic reagent (<i>M. musculus</i>)	B6.1239S4-Meox2tm1 (cre)Sor/J	Jackson Labs	Stock No 003755 RRID:IMSR_JAX:026858	Tallquist and Soriano, 2000 (another name <i>Meox2-Cre^{+/0}</i>)
Genetic reagent (<i>M. musculus</i>)	B6.129S-Snai1tm2Grid/J	Jackson Labs	Stock No 010686 RRID:IMSR_JAX:010686	Murray and Gridley, 2006 (another name <i>Snai1^{fllox/fllox}</i>)
Genetic reagent (<i>M. musculus</i>)	LuVeLu	Aulehla et al., 2008		Transgenic
Genetic reagent (<i>M. musculus</i>)	<i>T-str-Snai1</i>	This paper		Transgenic
Genetic reagent (<i>M. musculus</i>)	<i>Tgfr1^{+/-}</i>	This paper		Targeted null mutation
Genetic reagent (<i>M. musculus</i>)	<i>Cdx2-Alk5^{CA}</i>	Jurberg et al., 2013		Transgenic
Antibody	anti-Brachyury (Goat polyclonal)	R and D Systems	AF2085 RRID:AB_2200235	IF (1:200)
Antibody	Anti-Sox2 (Rabbit monoclonal)	Abcam	ab92494 RRID:AB_10585428	IF (1:200)
Antibody	anti-Cdh1 (Goat polyclonal)	R and D Systems	AF648 RRID:AB_355504	IF (1:200)
Antibody	anti-Cdh2 (Rabbit polyclonal)	Abcam	ab18203 RRID:AB_444317	IF (1:200)
Antibody	anti-Tbx6 (Goat polyclonal)	R and D Systems	AF4744 RRID:AB_2200834	IF (1:200)
Antibody	anti-Laminin 111 (Rabbit polyclonal)	Sigma	L9393 RRID:AB_477163	IF (1:200)
Antibody	EpCAM/TROP1 (Goat polyclonal)	R and D Systems	AF960 RRID:AB_355745	IF (1:200)
Antibody	anti-Vimentin (Rabbit monoclonal)	Abcam,	ab92547 RRID:AB_10562134	IF (1:200)
Antibody	anti-goat 488 (Donkey polyclonal)	Molecular Probes	A11055 RRID:AB_2534102	IF (1:1000)
Antibody	anti-rabbit 568 (Donkey polyclonal)	ThermoFisher Scientific	A10042 RRID:AB_2534017	IF (1:1000)
Recombinant DNA reagent	<i>T-str</i> promoter	Clements et al., 1996		Primitive streak-specific promoter of <i>T</i> (Brachyury)
Recombinant DNA reagent	<i>Snai1</i> cDNA	Nieto et al., 1992		
Sequenced-based reagent	Oligonucleotides	This paper		Table 1
Commercial assay, kit	Nextera XT index kit v2 Set B	Illumina	FC-131-2002	
Commercial assay, kit	Library Quant Kit	Illumina	LC480	

Continued on next page

Continued

Reagent type (species) or resource	Designation	Source or reference	Identifiers	Additional information
Commercial assay, kit	KAPA Library Quantification Kits	KAPA Biosystems	KK4854	
Software, algorithm	SC3	Kiselev et al., 2017	RRID:SCR_015953	
Software, algorithm	SPRING	Weinreb et al., 2018		

Mouse lines

Meox2-Cre^{+/-} (Tallquist and Soriano, 2000) and *Snai1^{flox/flox}* (Murray and Gridley, 2006) mouse strains were obtained from Jackson labs (B6.129S-*Snai1^{tm2Grid}/J* – Stock No 010686; B6.129S4-*Meox2^{tm1(cre)Sor}/J* – Stock No 003755). The LuVeLu reporter strain (Aulehla et al., 2008) was provided by Alexander Aulehla and Olivier Pourquié. *Snai1*-cKO embryos (*Meox2-Cre^{+/-}::Snai1^{flox/-}*) were obtained by crossing *Meox2-Cre^{+/-}::Snai1^{+/-}* males with *Snai1^{flox/flox}* females. *Meox2-Cre^{+/-}::Snai1^{flox/-}::LuVeLu^{+/-}* embryos were obtained from similar crosses but with the LuVeLu reporter introduced into *Snai1^{flox/flox}* genotype.

To obtain *T-str-Snai1* transgenic embryos, constructs were prepared by cloning a ~ 1,5 kb *Snai1* cDNA (Nieto et al., 1992) under the control of the PS-specific promoter of *T* (*Brachyury*) (Clements et al., 1996). These constructs were liberated from vector sequences, gel purified and used to produce transgenic embryos by pronuclear injection in FVB/N fertilized oocytes according to standard procedures (Hogan et al., 1994).

The *Tgfb1^{+/-}* line was generated by CRISPR/Cas9, inserting the TGATGATAGGATCC sequence, containing three stop codons and a *Bam*HI restriction site in frame with the open reading frame in exon 2. For this, a gRNA containing the targeting sequence TTGACCTAATTCCTCGAGAC was produced by in vitro transcription with the T7 promoter from a plasmid derived from the gRNA-basic (Casaca et al., 2016). The purified gRNA was microinjected into fertilized FVB/N mouse oocytes together with the Cas9 mRNA and the synthetic ssDNA 5'-ACCACAGACAAAGTTATACACAATAGTATGTGTATAGCTGAAATTGACCTAATTCCTCGATGATGATAGGATCCGACAGGCCATTTGTATGTGCACCATCTTCAAAAACAGGGGCAGTTACTACAACATATTGC-3', containing the stop codons flanked by 60 nucleotide-long homology arms. Genotyping of embryos was performed by PCR, on DNA obtained from yolk sacs or tail biopsies from embryos or mice, respectively, as previously described (Aires et al., 2019). The primers used for genotyping are specified in Table 1.

Ethical statement

Experiments involving animals carried out in the Oeiras laboratory followed the Portuguese (Portaria 1005/92) and European (Directive 2010/63/EU) legislations, concerning housing, husbandry, and welfare. The project was reviewed and approved by the Ethics Committee of 'Instituto Gulbenkian de Ciência' and by the Portuguese National Entity, 'Direcção Geral de Alimentação Veterinária' (license reference: 014308).

Single cell isolation

Wild type outbred MF1 mice were crossed to obtain early head fold embryos. Caudal lateral epiblast (CLE, comprising a region immediately lateral to the posterior edge of the node and lateral to the primitive streak, extending to about half the length of the primitive streak) (Wymeersch et al., 2016) was microdissected in M2 medium (Sigma, M7167), using hand-pulled solid glass needles. After removal of the majority underlying paraxial mesoderm, the different tissue pieces were pooled and after 5 min at 37°C in 0.05% trypsin/EDTA, they were placed in neutralization solution [10% fetal calf serum in phosphate-buffered saline (PBS)] and dissociated by pipetting into single cells. Single cells were then transferred to a tube containing 2% FCS in PBS, strained through the 35 µm mesh of a FACS tube (Corning, 352235) and 1 µg/ml of 4',6-Diamidino-2-Phenylindole, Dihydrochloride (DAPI; Thermo Fisher Scientific, D1306) was added. Single-cells were sorted by Fluorescence-activated cell sorting FACS) with a FACS Aria II (BD Biosciences) into individual wells of a 96-well PCR plate (BioRad, HSS9601) containing the ERCC RNA spike-in Mix (Ambion, #4456740) diluted (1:100.000) in a solution of 0,2% Triton-X100 (vol/vol) (Sigma, T9284), containing 2 U/ µl of RNase

Table 1. Primers used for genotyping.

<i>Snai1</i> deletion Fwd	CGGGCTTAGGTGTTTTTCAGAC
<i>Snai1</i> deletion Rev	TGAAAGCGGCTCTGTTCAGT
<i>Snai1^{fllox}</i> Fwd	TGAAAGCGGCTCTGTTCAGTG
<i>Snai1^{fllox}</i> Rev	CTGCTGCACCCCTACTATGTG
<i>Meox2-Cre</i> Fwd	CGAGTGATGAGGTTTCGAAG
<i>Meox2-Cre</i> Rev	CCTGATCCTGGCAATTTTCGGCT
LuVeLu Fwd	TGCTGCTGCCCCGACAACCACT
LuVeLu Rev	CTTGACAGCTCGTCCATGCC
<i>Snai1</i> transgenics Fwd	TTGTGTCTGCACGACCTGTGG
<i>Snai1</i> transgenics Rev	ATGGGGAGGTAGCAGGGTCAG
<i>Tgfb1</i> Fwd	TGTGAGACAGATGGTCTTTGC
<i>Tgfb1</i> mutant allele Rev	GGCCTGTCGGATCCTATCATC
<i>Tgfb1</i> WT allele Rev	ACATACAAATGGCCTGTCTCG

inhibitor (Takara Bio Europe, #2313A), 25% (v/v) of 100 μ M oligo dT₃₀VN (Biomers) and 25% (v/v) of dNTPs 10 mM (Invitrogen, #18427013) into RNase-free H₂O (Thermo Fisher Scientific, 11430615). Plates were sealed with Microseal F (Biorad, MSF1001), centrifuged at 4°C for 1 min at 2000 rpm and stored at –80°C.

Single-cell RNA sequencing

Full-length RNA-seq from sorted single cells was done using the Smart-seq2 method (*Picelli et al., 2014*). Libraries from a total of 91 single cells were prepared using a Nextera XT index kit v2 Set B (Illumina) and quantified both using Library Quant Kit (Illumina - LC480, KAPA Biosystems - KK4854) and the AATI Fragment Analyzer. Sequencing was done at the IGC Genomics facility (Illumina, Next-seq 500) at 5 million single end 75 bp reads per cell.

Single-cell RNA-seq analysis

RNA-seq analysis was done using R software. Sequences were submitted to the GEO repository, accession number GSE147100. Raw sequences were aligned to the GRCm38 (mm10) reference genome using Hisat2 (*Kim et al., 2013*). The ‘featureCounts’ R function was used to count reads mapping to annotated genes. Clustering analysis was done using the single-cell consensus clustering (SC3) pipeline (*Kiselev et al., 2017*). Gene markers were obtained considering p-value<0,05. P-values are shown in Spreadsheet 1. Single-cell data visualization was done using SPRING (*Weinreb et al., 2018*), with the following parameters: Minimum UMI total (for filtering cells)=1000; Minimum number of cells with ≥ 3 counts (for filtering genes) was set to 3; Gene variability percentile (for filtering genes)=50; Number of PCA dimensions (for building graph) was set to 20; Number of nearest neighbours (for graph)=5. Single cell heatmaps were created using the Heatmapper platform (*Babicki et al., 2016*). The scRNA-seq values in RPKM used for this analysis are shown Spreadsheet 2.

In Situ hybridization and sectioning

Wholemount in situ hybridization was performed as previously described (*Aires et al., 2019*). These experiments were repeated independently at least twice for each genotype giving identical results, with the exception of *Lfng*, which gave different patterns in the PSM region as reported in **Figure 5** and **Figure 5—figure supplement 1**, and notochord markers in *Snai1*-cKO embryos, that gave two alternative patterns as referenced in the main text. Post-stained embryos were included in a mixture of 0.45% gelatin (Merck), 27% bovine serum albumin (Roche), 18% sucrose (Sigma) in PBS that was then jellified with 1.75% glutaraldehyde (Biochem chemopharma) and sectioned at 20 μ m with a vibratome (Leica).

Immunohistochemistry and 3D imaging

Wholemount immunofluorescence staining of tail tissues was performed as previously described (Osorno *et al.*, 2012). At least two embryos per genotype were stained with each antibody with similar results. Primary antibodies (1:200): goat anti-Brachyury (R and D Systems, AF2085), rabbit anti-Sox2 (Abcam, AB92494), goat anti-Cdh1 (R and D Systems, AF648), rabbit anti-Cdh2 (Abcam, AB18203), goat anti-Tbx6 (R and D Systems, AF4744), rabbit anti-Laminin 111 which detects all laminins containing $\alpha 1$, $\beta 1$ or $\gamma 1$ chains (Sigma, L9393), goat anti-human EpCAM/TROP1 (R and D Systems, AF960) and rabbit anti-Vimentin (Abcam, AB92547). Secondary antibodies (1:1000): donkey anti-goat 488 (Molecular Probes, A-11055) and donkey anti-rabbit 568 (Molecular Probes, A10042). Immuno-stained tails were imaged on a Prairie two-photon system, using an Olympus 20 \times 1.0 NA W objective, with the excitation laser tuned to 960 nm, and GaAsP photodetectors. Z stacks of 1024 \times 1024 images were acquired every 1 μ m, with either 1x or 1.5x zoom. Laser intensity and photomultiplier levels were maintained across replicates and controls.

Live imaging (LuVeLu)

Embryos expressing the LuVeLu reporter (Aulehla *et al.*, 2008) were dissected in pre-warmed (at 37°C) M2 medium (Sigma) and cultured in low glucose DMEM medium (Gibco, 11054020), 10% of HyClone defined fetal bovine serum (GE Healthcare, #HYCLSH30070.03), 2 mM of L-glutamine (Gibco, #25030-024) and 1% penicillin-streptomycin (Sigma, #P0781). Embryos were cultured at 37°C in a 65% O₂ and 15% CO₂ environment (N2 balanced). Embryos were imaged on the Prairie two-photon system (laser tuned to 960 nm). At time = 0, a z-stack of 5 μ m step-size was acquired at 1024 \times 1024 pixel size, using the 20x objective. From t = 1 and onwards, we acquired z-stacks series of 1024 \times 512 pixel images, spaced in depth at 10 μ m, using a Nikon 16x LWD 0.8NA W objective. T-series were acquired every 8,5 min.

Image processing

3D and 4D series of two-photon microscopy datasets were processed using Fiji (Schindelin *et al.*, 2012). Pre-processing involved removal of outlier pixels and elimination of electronic noise in the form of periodic patterns by using fast-Fourier filters (detection of 'maxima' in the Fourier spectrum and deleting for each an area with $r = 5$ pixels, performed slice-by-slice). 3D datasets of immuno-stained tissues were deconvolved using Huygens (SVI). When acquired, adjacent 3-stack datasets were digitally stitched using the 'Image Stitching' plugin (Preibisch *et al.*, 2009) (pairwise mode, with alpha blending). We then evaluated the Z-axis attenuation in 3D datasets; when attenuation in depth was noticeable we performed a compensation using FIJI's 'Math...Batch' function and the expression: $v = v * \exp(a * z)$, where v = pixel intensity; a = compensation factor (ranging from 1 to 2, depending on the compensation necessary to equalize the intensities on the first and the deeper optical slices) – replicates and controls were treated similarly. The pre-processed multi-channel dataset was then repositioned by affine transformation using the 'TransformJ' plugin (Meijering *et al.*, 2001). The transformation matrix was obtained using FIJI's 3D viewer plugin. 3D visualization and analysis of embryo tails was done using Imaris v9.0 (Bitplane), including the rendering of wholemount (blend mode) and of sagittal and transversal 5 μ m optical sections by maximum intensity projection. Note: In EPCAM-stained samples, a considerable bleed-through from the DAPI channel was observed. Therefore, before deconvolution, the EPCAM channel was compensated by dividing it by the DAPI signal. Further background reduction was achieved by dividing the compensated dataset by a 'synthetic background' obtained from optical slices without stained tissue. Mutant and wild type embryos were treated similarly.

A CLAHE filter was used to enhance the contrast of DAPI-stained embryos which were also imaged using the Prairie two-photon system (20x objective and laser tuned to 890 nm). Defined embryonic structures (e.g. neural tube) were then segmented by manual contouring and 3D rendered using Amira (Thermo-Fisher Scientific).

To analyse the LuVeLu time-series, we concatenated all 3D datasets and converted to a 4D dataset in HDF5 format. The time-points were then registered using the BigStitcher plugin (Hörl *et al.*, 2019). Visualization and analysis were done using Imaris.

Acknowledgements

We would like to thank Achim Gossler for the *T-Str* enhancer, Olivier Pourquié and Alexander Aulehla for the LuVeLu reporter strain, Sólveig Thorsteinsdóttir for the laminin antibody, Angela Nieto for *Snai1* cDNA, Julien Delile for sharing data files, Luisa de Lemos for sharing unpublished data, Daniel Neves for the help with the analysis of the RNA-seq data, Hugo Pereira for the help using BigStitcher and Nuno Granjeiro for helping to set up the live imaging apparatus. We also would like to thank the IGC and SCRM animal facilities, Fiona Rossi and Clair Cryer from the SCRM flow cytometry facility, João Sobral from the IGC genomics facility, and past and present members of the Mallo lab for useful comments and support during the course of this project. This work has been supported by grants PTDC/BEX-BID/0899/2014 and LISBOA-01-0145-FEDER-030254 (FCT, Portugal) and SCML-MC-60-2014 (Santa Casa da Misericórdia, Portugal) to M.M.; the research infrastructure Congento, project LISBOA-01-0145-FEDER-022170; PhD fellowships PD/BD/128426/2017 to AD and (PD/BD/128437/2017) to AL (FCT, Portugal); MRC (UK) grants MR/S008799/1 and MR/K011200/1 to V.W. and a Development Travelling Fellowship (DEV-170806) from 'The Company of Biologists' to AD.

Additional information

Funding

Funder	Grant reference number	Author
Fundação para a Ciência e a Tecnologia	PTDC/BEX-BID/0899/2014	Moises Mallo
Fundação para a Ciência e a Tecnologia	LISBOA-01-0145-FEDER-030254	Moises Mallo
Fundação para a Ciência e a Tecnologia	LISBOA-01-0145-FEDER-022170	Moises Mallo
Fundação para a Ciência e a Tecnologia	PD/BD/128426/2017	André Dias
Fundação para a Ciência e a Tecnologia	PD/BD/128437/2017	Anastasiia Lozovska
Santa Casa da Misericórdia	SCML-MC-60-2014	Moises Mallo
Medical Research Council	MR/S008799/1	Valerie Wilson
Medical Research Council	MR/K011200/1	Valerie Wilson
Company of Biologists	DEV-170806	André Dias

The funders had no role in study design, data collection and interpretation, or the decision to submit the work for publication.

Author contributions

André Dias, Conceptualization, Formal analysis, Investigation, Writing - review and editing; Anastasiia Lozovska, Ana Nóvoa, Investigation; Filip J Wymeersch, Anahi Binagui-Casas, Investigation, Methodology; Daniel Sobral, Formal analysis, Methodology; Gabriel G Martins, Data curation, Methodology; Valerie Wilson, Supervision, Funding acquisition, Methodology; Moises Mallo, Conceptualization, Supervision, Funding acquisition, Investigation, Writing - original draft, Project administration, Writing - review and editing

Author ORCIDs

André Dias  <https://orcid.org/0000-0003-3337-6373>

Anastasiia Lozovska  <https://orcid.org/0000-0002-9842-6450>

Filip J Wymeersch  <https://orcid.org/0000-0001-8999-4555>

Ana Nóvoa  <https://orcid.org/0000-0002-5668-5630>

Anahi Binagui-Casas  <https://orcid.org/0000-0002-7987-9286>

Daniel Sobral  <https://orcid.org/0000-0003-3955-0117>

Gabriel G Martins  <http://orcid.org/0000-0002-6506-9776>

Valerie Wilson  <https://orcid.org/0000-0003-4182-5159>

Moises Mallo  <https://orcid.org/0000-0002-9744-0912>

Ethics

Animal experimentation: Experiments involving animals carried out in the Oeiras laboratory followed the Portuguese (Portaria 1005/92) and European (Directive 2010/63/EU) legislations, concerning housing, husbandry, and welfare. The project was reviewed and approved by the Ethics Committee of "Instituto Gulbenkian de Ciência" and by the Portuguese National Entity, "Direcção Geral de Alimentação Veterinária" (license reference: 014308).

Decision letter and Author response

Decision letter <https://doi.org/10.7554/eLife.56615.sa1>

Author response <https://doi.org/10.7554/eLife.56615.sa2>

Additional files

Supplementary files

- Transparent reporting form

Data availability

Sequencing data have been deposited in GEO under accession code GSE147100.

The following dataset was generated:

Author(s)	Year	Dataset title	Dataset URL	Database and Identifier
Dias A, Mallo M	2020	Single-cell RNA sequencing of neuromesodermal progenitors of early headfold wild type embryos	https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE147100	NCBI Gene Expression Omnibus, GSE147100

The following previously published datasets were used:

Author(s)	Year	Dataset title	Dataset URL	Database and Identifier
Aires R, Mallo M	2018	RNA-Seq of mus musculus: Tailbud WT1	https://www.ncbi.nlm.nih.gov/sra/SRX4968732	NCBI Sequence Read Archive, SRX4968732
Aires R, Mallo M	2018	RNA-Seq of mus musculus: Tailbud WT2	https://www.ncbi.nlm.nih.gov/sra/SRX4968731	NCBI Sequence Read Archive, SRX4968731
deLemos L, Mallo M	2019	RNA-seq of mus musculus : Tail Bud progenitors 2	https://www.ncbi.nlm.nih.gov/sra/SRX5532193	NCBI Sequence Read Archive, SRX5532193
deLemos L, Mallo M	2019	RNA-seq of mus musculus : Tail Bud progenitors 1	https://www.ncbi.nlm.nih.gov/sra/SRX5532192	NCBI Sequence Read Archive, SRX5532192
Wymeersch FJ, Skylaki S, Huang Y, Watson JA, Economou C, Marek-Johnston C, Tomlinson SR, Wilson V	2018	Gene expression in microdissected embryonic regions during mouse axis elongation.	https://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE120870	NCBI Gene Expression Omnibus, GSE120870
Briscoe J, Kleijung J, Delile J, Gouti M	2015	Retinoic acid mediated neural and mesoderm specification during vertebrate trunk	https://www.ebi.ac.uk/arrayexpress/experiments/E-MTAB-5208	ArrayExpress, E-MTAB-5208

References

- Acloque H**, Adams MS, Fishwick K, Bronner-Fraser M, Nieto MA. 2009. Epithelial-mesenchymal transitions: the importance of changing cell state in development and disease. *Journal of Clinical Investigation* **119**:1438–1449. DOI: <https://doi.org/10.1172/JCI38019>, PMID: 19487820
- Aires R**, Jurberg AD, Leal F, Nóvoa A, Cohn MJ, Mallo M. 2016. Oct4 is a key regulator of vertebrate trunk length diversity. *Developmental Cell* **38**:262–274. DOI: <https://doi.org/10.1016/j.devcel.2016.06.021>, PMID: 27453501
- Aires R**, Dias A, Mallo M. 2018. Deconstructing the molecular mechanisms shaping the vertebrate body plan. *Current Opinion in Cell Biology* **55**:81–86. DOI: <https://doi.org/10.1016/j.ceb.2018.05.009>, PMID: 30015151
- Aires R**, de Lemos L, Nóvoa A, Jurberg AD, Mascres B, Duboule D, Mallo M. 2019. Tail bud progenitor activity relies on a network comprising Gdf11, Lin28, and Hox13 genes. *Developmental Cell* **48**:383–395. DOI: <https://doi.org/10.1016/j.devcel.2018.12.004>, PMID: 30661984
- Anderson MJ**, Naiche LA, Wilson CP, Elder C, Swing DA, Lewandoski M. 2013. TCreERT2, a transgenic mouse line for temporal control of Cre-mediated recombination in lineages emerging from the primitive streak or tail bud. *PLOS ONE* **8**:e62479. DOI: <https://doi.org/10.1371/journal.pone.0062479>, PMID: 23638095
- Andersson O**, Reissmann E, Ibáñez CF. 2006. Growth differentiation factor 11 signals through the transforming growth factor-beta receptor ALK5 to regionalize the anterior-posterior Axis. *EMBO Reports* **7**:831–837. DOI: <https://doi.org/10.1038/sj.embor.7400752>, PMID: 16845371
- Andre P**, Song H, Kim W, Kispert A, Yang Y. 2015. Wnt5a and Wnt11 regulate mammalian anterior-posterior Axis elongation. *Development* **142**:1516–1527. DOI: <https://doi.org/10.1242/dev.119065>, PMID: 25813538
- Arnold SJ**, Hofmann UK, Bikoff EK, Robertson EJ. 2008. Pivotal roles for eomesodermin during Axis formation, epithelium-to-mesenchyme transition and endoderm specification in the mouse. *Development* **135**:501–511. DOI: <https://doi.org/10.1242/dev.014357>, PMID: 18171685
- Aulehla A**, Wiegand W, Baubet V, Wahl MB, Deng C, Taketo M, Lewandoski M, Pourquié O. 2008. A beta-catenin gradient links the clock and wavefront systems in mouse embryo segmentation. *Nature Cell Biology* **10**:186–193. DOI: <https://doi.org/10.1038/ncb1679>, PMID: 18157121
- Babicki S**, Arndt D, Marcu A, Liang Y, Grant JR, Maciejewski A, Wishart DS. 2016. Heatmapper: web-enabled heat mapping for all. *Nucleic Acids Research* **44**:W147–W153. DOI: <https://doi.org/10.1093/nar/gkw419>, PMID: 27190236
- Barrallo-Gimeno A**, Nieto MA. 2005. The snail genes as inducers of cell movement and survival: implications in development and Cancer. *Development* **132**:3151–3161. DOI: <https://doi.org/10.1242/dev.01907>, PMID: 15983400
- Bénazéraf B**, Pourquié O. 2013. Formation and segmentation of the vertebrate body Axis. *Annual Review of Cell and Developmental Biology* **29**:1–26. DOI: <https://doi.org/10.1146/annurev-cellbio-101011-155703>, PMID: 23808844
- Boulet AM**, Capecchi MR. 2012. Signaling by FGF4 and FGF8 is required for axial elongation of the mouse embryo. *Developmental Biology* **371**:235–245. DOI: <https://doi.org/10.1016/j.ydbio.2012.08.017>, PMID: 22954964
- Cambray N**, Wilson V. 2002. Axial progenitors with extensive potency are localised to the mouse chordoneural hinge. *Development* **129**:4855–4866. PMID: 12361976
- Cambray N**, Wilson V. 2007. Two distinct sources for a population of maturing axial progenitors. *Development* **134**:2829–2840. DOI: <https://doi.org/10.1242/dev.02877>, PMID: 17611225
- Carver EA**, Jiang R, Lan Y, Oram KF, Gridley T. 2001. The mouse snail gene encodes a key regulator of the epithelial-mesenchymal transition. *Molecular and Cellular Biology* **21**:8184–8188. DOI: <https://doi.org/10.1128/MCB.21.23.8184-8188.2001>, PMID: 11689706
- Casaca A**, Nóvoa A, Mallo M. 2016. Hoxb6 can interfere with somitogenesis in the posterior embryo through a mechanism independent of its rib-promoting activity. *Development* **143**:437–448. DOI: <https://doi.org/10.1242/dev.133074>, PMID: 26718008
- Catala M**, Teillet MA, Le Douarin NM. 1995. Organization and development of the tail bud analyzed with the quail-chick Chimaera system. *Mechanisms of Development* **51**:51–65. DOI: [https://doi.org/10.1016/0925-4773\(95\)00350-A](https://doi.org/10.1016/0925-4773(95)00350-A), PMID: 7669693
- Clements D**, Taylor HC, Herrmann BG, Stott D. 1996. Distinct regulatory control of the brachyury gene in axial and non-axial mesoderm suggests separation of mesoderm lineages early in mouse gastrulation. *Mechanisms of Development* **56**:139–149. DOI: [https://doi.org/10.1016/0925-4773\(96\)00520-5](https://doi.org/10.1016/0925-4773(96)00520-5), PMID: 8798154
- de Lemos L**, Nóvoa A, Mallo M. 2019. High EphA1 expression is a potential cell surface marker for embryonic neuro-mesodermal progenitors. *bioRxiv*. DOI: <https://doi.org/10.1101/584524>
- Deng CX**, Wynshaw-Boris A, Shen MM, Daugherty C, Ornitz DM, Leder P. 1994. Murine FGFR-1 is required for early postimplantation growth and axial organization. *Genes & Development* **8**:3045–3057. DOI: <https://doi.org/10.1101/gad.8.24.3045>, PMID: 8001823
- Derynck R**, Muthusamy BP, Saetern KY. 2014. Signaling pathway cooperation in TGF- β -induced epithelial-mesenchymal transition. *Current Opinion in Cell Biology* **31**:56–66. DOI: <https://doi.org/10.1016/j.ceb.2014.09.001>, PMID: 25240174
- DeVeale B**, Brokman I, Mohseni P, Babak T, Yoon C, Lin A, Onishi K, Tomilin A, Pevny L, Zandstra PW, Nagy A, van der Kooy D. 2013. Oct4 is required ~E7.5 for proliferation in the primitive streak. *PLOS Genetics* **9**:e1003957. DOI: <https://doi.org/10.1371/journal.pgen.1003957>, PMID: 24244203

- Gouti M**, Delile J, Stamatakis D, Wymeersch FJ, Huang Y, Kleinjung J, Wilson V, Briscoe J. 2017. A gene regulatory network balances neural and mesoderm specification during vertebrate trunk development. *Developmental Cell* **41**:243–261. DOI: <https://doi.org/10.1016/j.devcel.2017.04.002>, PMID: 28457792
- Greco TL**, Takada S, Newhouse MM, McMahon JA, McMahon AP, Camper SA. 1996. Analysis of the vestigial tail mutation demonstrates that Wnt-3a gene dosage regulates mouse axial development. *Genes & Development* **10**:313–324. DOI: <https://doi.org/10.1101/gad.10.3.313>, PMID: 8595882
- Henrique D**, Abranches E, Verrier L, Storey KG. 2015. Neuromesodermal progenitors and the making of the spinal cord. *Development* **142**:2864–2875. DOI: <https://doi.org/10.1242/dev.119768>, PMID: 26329597
- Hernández-Martínez R**, Ramkumar N, Anderson KV. 2019. p120-catenin regulates WNT signaling and EMT in the mouse embryo. *PNAS* **116**:16872–16881. DOI: <https://doi.org/10.1073/pnas.1902843116>, PMID: 31371508
- Herrmann BG**, Labeit S, Poustka A, King TR, Lehrach H. 1990. Cloning of the T gene required in mesoderm formation in the mouse. *Nature* **343**:617–622. DOI: <https://doi.org/10.1038/343617a0>, PMID: 2154694
- Hogan B**, Beddington R, Constantini F, Lacy E. 1994. *Manipulating the Mouse Embryo: A Laboratory Manual*. Cold Spring Harbour Laboratory Press.
- Holmdahl DE**. 1925. Experimentelle untersuchungen über die lage der grenze primärer und sekundärer körperentwicklung beim huhn. *Anatomischer Anzeiger* **59**:393–396.
- Hörl D**, Rojas Rusak F, Preusser F, Tillberg P, Randel N, Chhetri RK, Cardona A, Keller PJ, Harz H, Leonhardt H, Treier M, Preibisch S. 2019. BigStitcher: reconstructing high-resolution image datasets of cleared and expanded samples. *Nature Methods* **16**:870–874. DOI: <https://doi.org/10.1038/s41592-019-0501-0>, PMID: 31384047
- Javali A**, Misra A, Leonavicius K, Acharyya D, Vyas B, Sambasivan R. 2017. Co-expression of Tbx6 and Sox2 identifies a novel transient neuromesoderm progenitor cell state. *Development* **144**:4522–4529. DOI: <https://doi.org/10.1242/dev.153262>, PMID: 29084802
- Jurberg AD**, Aires R, Varela-Lasheras I, Nóvoa A, Mallo M. 2013. Switching axial progenitors from producing trunk to tail tissues in vertebrate embryos. *Developmental Cell* **25**:451–462. DOI: <https://doi.org/10.1016/j.devcel.2013.05.009>, PMID: 23763947
- Kalluri R**, Weinberg RA. 2009. The basics of epithelial-mesenchymal transition. *Journal of Clinical Investigation* **119**:1420–1428. DOI: <https://doi.org/10.1172/JCI39104>, PMID: 19487818
- Kim D**, Pertea G, Trapnell C, Pimentel H, Kelley R, Salzberg SL. 2013. TopHat2: accurate alignment of transcriptomes in the presence of insertions, deletions and gene fusions. *Genome Biology* **14**:R36. DOI: <https://doi.org/10.1186/gb-2013-14-4-r36>, PMID: 23618408
- Kiselev VY**, Kirschner K, Schaub MT, Andrews T, Yiu A, Chandra T, Natarajan KN, Reik W, Barahona M, Green AR, Hemberg M. 2017. SC3: consensus clustering of single-cell RNA-seq data. *Nature Methods* **14**:483–486. DOI: <https://doi.org/10.1038/nmeth.4236>, PMID: 28346451
- Koch F**, Scholze M, Wittler L, Schifferl D, Sudheer S, Grote P, Timmermann B, Macura K, Herrmann BG. 2017. Antagonistic activities of Sox2 and brachyury control the fate choice of Neuro-Mesodermal progenitors. *Developmental Cell* **42**:514–526. DOI: <https://doi.org/10.1016/j.devcel.2017.07.021>, PMID: 28826820
- Lamouille S**, Xu J, Derynck R. 2014. Molecular mechanisms of epithelial-mesenchymal transition. *Nature Reviews Molecular Cell Biology* **15**:178–196. DOI: <https://doi.org/10.1038/nrm3758>, PMID: 24556840
- Liu P**, Wakamiya M, Shea MJ, Albrecht U, Behringer RR, Bradley A. 1999. Requirement for Wnt3 in vertebrate Axis formation. *Nature Genetics* **22**:361–365. DOI: <https://doi.org/10.1038/11932>, PMID: 10431240
- Liu JP**. 2006. The function of growth/differentiation factor 11 (Gdf11) in Rostrocaudal patterning of the developing spinal cord. *Development* **133**:2865–2874. DOI: <https://doi.org/10.1242/dev.02478>, PMID: 16790475
- Lomeli H**, Starling C, Gridley T. 2009. Epiblast-specific Snai1 deletion results in embryonic lethality due to multiple vascular defects. *BMC Research Notes* **2**:22. DOI: <https://doi.org/10.1186/1756-0500-2-22>, PMID: 19284699
- Mallo M**. 2018. Reassessing the role of hox genes during vertebrate development and evolution. *Trends in Genetics* **34**:209–217. DOI: <https://doi.org/10.1016/j.tig.2017.11.007>
- Martin BL**, Kimelman D. 2012. Canonical wnt signaling dynamically controls multiple stem cell fate decisions during vertebrate body formation. *Developmental Cell* **22**:223–232. DOI: <https://doi.org/10.1016/j.devcel.2011.11.001>, PMID: 22264734
- Matsubara Y**, Hirasawa T, Egawa S, Hattori A, Suganuma T, Kohara Y, Nagai T, Tamura K, Kuratani S, Kuroiwa A, Suzuki T. 2017. Anatomical integration of the sacral-hindlimb unit coordinated by GDF11 underlies variation in hindlimb positioning in tetrapods. *Nature Ecology & Evolution* **1**:1392–1399. DOI: <https://doi.org/10.1038/s41559-017-0247-y>, PMID: 29046533
- McPherron AC**, Lawler AM, Lee SJ. 1999. Regulation of anterior/posterior patterning of the axial skeleton by growth/differentiation factor 11. *Nature Genetics* **22**:260–264. DOI: <https://doi.org/10.1038/10320>, PMID: 10391213
- McPherron AC**, Huynh TV, Lee SJ. 2009. Redundancy of myostatin and growth/differentiation factor 11 function. *BMC Developmental Biology* **9**:24. DOI: <https://doi.org/10.1186/1471-213X-9-24>, PMID: 19298661
- Meijering EH**, Niessen WJ, Viergever MA. 2001. Quantitative evaluation of convolution-based methods for medical image interpolation. *Medical Image Analysis* **5**:111–126. DOI: [https://doi.org/10.1016/S1361-8415\(00\)00040-2](https://doi.org/10.1016/S1361-8415(00)00040-2), PMID: 11516706
- Murray SA**, Gridley T. 2006. Snail family genes are required for left-right asymmetry determination, but not neural crest formation, in mice. *PNAS* **103**:10300–10304. DOI: <https://doi.org/10.1073/pnas.0602234103>, PMID: 16801545

- Naiche LA**, Holder N, Lewandoski M. 2011. FGF4 and FGF8 comprise the wavefront activity that controls somitogenesis. *PNAS* **108**:4018–4023. DOI: <https://doi.org/10.1073/pnas.1007417108>, PMID: 21368122
- Nieto MA**, Bennett MF, Sargent MG, Wilkinson DG. 1992. Cloning and developmental expression of *sna*, a murine homologue of the *Drosophila* *snail* gene. *Development* **116**:227–237. PMID: 1483390
- Nieto MA**. 2002. The snail superfamily of zinc-finger transcription factors. *Nature Reviews Molecular Cell Biology* **3**:155–166. DOI: <https://doi.org/10.1038/nrm757>, PMID: 11994736
- Nieto MA**, Huang RY, Jackson RA, Thiery JP. 2016. EMT. 2016. *Cell* **166**:21–45. DOI: <https://doi.org/10.1016/j.cell.2016.06.028>, PMID: 27368099
- Olivera-Martinez I**, Harada H, Halley PA, Storey KG. 2012. Loss of FGF-dependent mesoderm identity and rise of endogenous retinoid signalling determine cessation of body Axis elongation. *PLOS Biology* **10**:e1001415. DOI: <https://doi.org/10.1371/journal.pbio.1001415>, PMID: 23118616
- Osorno R**, Tsakiridis A, Wong F, Cambay N, Economou C, Wilkie R, Blin G, Scotting PJ, Chambers I, Wilson V. 2012. The developmental dismantling of pluripotency is reversed by ectopic Oct4 expression. *Development* **139**:2288–2298. DOI: <https://doi.org/10.1242/dev.078071>, PMID: 22669820
- Pastushenko I**, Brisebarre A, Sifrim A, Fioramonti M, Revenco T, Boumahdi S, Van Keymeulen A, Brown D, Moers V, Lemaire S, De Clercq S, Minguijón E, Balsat C, Sokolow Y, Dubois C, De Cock F, Scozzaro S, Sopena F, Lanas A, D'Haene N, et al. 2018. Identification of the tumour transition states occurring during EMT. *Nature* **556**:463–468. DOI: <https://doi.org/10.1038/s41586-018-0040-3>, PMID: 29670281
- Picelli S**, Faridani OR, Björklund AK, Winberg G, Sagasser S, Sandberg R. 2014. Full-length RNA-seq from single cells using Smart-seq2. *Nature Protocols* **9**:171–181. DOI: <https://doi.org/10.1038/nprot.2014.006>, PMID: 24385147
- Preibisch S**, Saalfeld S, Tomancak P. 2009. Globally optimal stitching of tiled 3D microscopic image acquisitions. *Bioinformatics* **25**:1463–1465. DOI: <https://doi.org/10.1093/bioinformatics/btp184>, PMID: 19346324
- Ramkumar N**, Omelchenko T, Silva-Gagliardi NF, McGlade CJ, Wijnholds J, Anderson KV. 2016. Crumbs2 promotes cell ingression during the epithelial-to-mesenchymal transition at Gastrulation. *Nature Cell Biology* **18**:1281–1291. DOI: <https://doi.org/10.1038/ncb3442>, PMID: 27870829
- Robinton DA**, Chal J, Lummertz da Rocha E, Han A, Yermalovich AV, Oginuma M, Schlaeger TM, Sousa P, Rodriguez A, Urbach A, Pourquié O, Daley GQ. 2019. The Lin28/let-7 pathway regulates the mammalian caudal body Axis elongation program. *Developmental Cell* **48**:396–405. DOI: <https://doi.org/10.1016/j.devcel.2018.12.016>, PMID: 30661985
- Rodrigo Albors A**, Halley PA, Storey KG. 2018. Lineage tracing of axial progenitors using Nkx1-2CreER^{T2} mice defines their trunk and tail contributions. *Development* **145**:dev164319. DOI: <https://doi.org/10.1242/dev.164319>, PMID: 30201686
- Saitsu H**, Yamada S, Uwabe C, Ishibashi M, Shiota K. 2004. Development of the posterior neural tube in human embryos. *Anatomy and Embryology* **209**:107–117. DOI: <https://doi.org/10.1007/s00429-004-0421-2>, PMID: 15597189
- Saitsu H**, Shiota K. 2008. Involvement of the axially condensed tail bud mesenchyme in normal and abnormal human posterior neural tube development. *Congenital Anomalies* **48**:1–6. DOI: <https://doi.org/10.1111/j.1741-4520.2007.00178.x>, PMID: 18230116
- Savory JG**, Mansfield M, Rijli FM, Lohnes D. 2011. Cdx mediates neural tube closure through transcriptional regulation of the planar cell polarity gene *Ptk7*. *Development* **138**:1361–1370. DOI: <https://doi.org/10.1242/dev.056622>, PMID: 21350009
- Schindelin J**, Arganda-Carreras I, Frise E, Kaynig V, Longair M, Pietzsch T, Preibisch S, Rueden C, Saalfeld S, Schmid B, Tinevez JY, White DJ, Hartenstein V, Eliceiri K, Tomancak P, Cardona A. 2012. Fiji: an open-source platform for biological-image analysis. *Nature Methods* **9**:676–682. DOI: <https://doi.org/10.1038/nmeth.2019>, PMID: 22743772
- Schoenwolf GC**. 1984. Histological and ultrastructural studies of secondary neurulation in mouse embryos. *American Journal of Anatomy* **169**:361–376. DOI: <https://doi.org/10.1002/aja.1001690402>, PMID: 6731331
- Schoenwolf GC**, Smith JL. 1990. Mechanisms of neurulation: traditional viewpoint and recent advances. *Development* **109**:243–270. PMID: 2205465
- Shifley ET**, Vanhorn KM, Perez-Balaguer A, Franklin JD, Weinstein M, Cole SE. 2008. Oscillatory lunatic fringe activity is crucial for segmentation of the anterior but not posterior skeleton. *Development* **135**:899–908. DOI: <https://doi.org/10.1242/dev.006742>, PMID: 18234727
- Stern CD**, Charité J, Deschamps J, Duboule D, Durston AJ, Kmita M, Nicolas JF, Palmeirim I, Smith JC, Wolpert L. 2006. Head-tail patterning of the vertebrate embryo: one, two or many unresolved problems? *The International Journal of Developmental Biology* **50**:3–15. DOI: <https://doi.org/10.1387/ijdb.052095cs>, PMID: 16323073
- Steventon B**, Martinez Arias A. 2017. Evo-engineering and the cellular and molecular origins of the vertebrate spinal cord. *Developmental Biology* **432**:3–13. DOI: <https://doi.org/10.1016/j.ydbio.2017.01.021>, PMID: 28192080
- Takada S**, Stark KL, Shea MJ, Vassileva G, McMahon JA, McMahon AP. 1994. Wnt-3a regulates somite and tailbud formation in the mouse embryo. *Genes & Development* **8**:174–189. DOI: <https://doi.org/10.1101/gad.8.2.174>, PMID: 8299937
- Tallquist MD**, Soriano P. 2000. Epiblast-restricted cre expression in MORE mice: a tool to distinguish embryonic vs. extra-embryonic gene function. *Genesis* **26**:113–115. DOI: [https://doi.org/10.1002/\(SICI\)1526-968X\(200002\)26:2<113::AID-GENE3>3.0.CO;2-2](https://doi.org/10.1002/(SICI)1526-968X(200002)26:2<113::AID-GENE3>3.0.CO;2-2), PMID: 10686601

- Tam PP, Behringer RR. 1997. Mouse gastrulation: the formation of the mammalian body plan. *Mechanisms of Development* **68**:3–25. DOI: [https://doi.org/10.1016/s0925-4773\(97\)00123-8](https://doi.org/10.1016/s0925-4773(97)00123-8), PMID: 9431800
- Tsakiridis A, Huang Y, Blin G, Skylaki S, Wymeersch F, Osorno R, Economou C, Karagianni E, Zhao S, Lowell S, Wilson V. 2014. Distinct Wnt-driven primitive streak-like populations reflect in vivo lineage precursors. *Development* **141**:1209–1221. DOI: <https://doi.org/10.1242/dev.101014>, PMID: 24595287
- Tzouanacou E, Wegener A, Wymeersch FJ, Wilson V, Nicolas JF. 2009. Redefining the progression of lineage segregations during mammalian embryogenesis by clonal analysis. *Developmental Cell* **17**:365–376. DOI: <https://doi.org/10.1016/j.devcel.2009.08.002>, PMID: 19758561
- Voiculescu O, Bodenstein L, Lau I-J, Stern CD. 2014. Local cell interactions and self-amplifying individual cell ingression drive amniote gastrulation. *eLife* **3**:e01817. DOI: <https://doi.org/10.7554/eLife.01817>
- Weinreb C, Wolock S, Klein AM. 2018. SPRING: a kinetic interface for visualizing high dimensional single-cell expression data. *Bioinformatics* **34**:1246–1248. DOI: <https://doi.org/10.1093/bioinformatics/btx792>
- Williams DR, Shifley ET, Lather JD, Cole SE. 2014. Posterior skeletal development and the segmentation clock period are sensitive to *lfrng* dosage during somitogenesis. *Developmental Biology* **388**:159–169. DOI: <https://doi.org/10.1016/j.ydbio.2014.02.006>, PMID: 24560643
- Wilson V, Olivera-Martinez I, Storey KG. 2009. Stem cells, signals and vertebrate body Axis extension. *Development* **136**:2133. DOI: <https://doi.org/10.1242/dev.039172>
- Wymeersch FJ, Huang Y, Blin G, Cambay N, Wilkie R, Wong FC, Wilson V. 2016. Position-dependent plasticity of distinct progenitor types in the primitive streak. *eLife* **5**:e10042. DOI: <https://doi.org/10.7554/eLife.10042>, PMID: 26780186
- Wymeersch FJ, Skylaki S, Huang Y, Watson JA, Economou C, Marek-Johnston C, Tomlinson SR, Wilson V. 2019. Transcriptionally dynamic progenitor populations organised around a stable niche drive axial patterning. *Development* **146**:dev168161. DOI: <https://doi.org/10.1242/dev.168161>, PMID: 30559277
- Xu J, Lamouille S, Derynck R. 2009. TGF-beta-induced epithelial to mesenchymal transition. *Cell Research* **19**:156–172. DOI: <https://doi.org/10.1038/cr.2009.5>, PMID: 19153598
- Yamaguchi TP, Harpal K, Henkemeyer M, Rossant J. 1994. *fgfr-1* is required for embryonic growth and mesodermal patterning during mouse gastrulation. *Genes & Development* **8**:3032–3044. DOI: <https://doi.org/10.1101/gad.8.24.3032>, PMID: 8001822
- Zeisberg M, Neilson EG. 2009. Biomarkers for epithelial-mesenchymal transitions. *Journal of Clinical Investigation* **119**:1429–1437. DOI: <https://doi.org/10.1172/JCI36183>, PMID: 19487819